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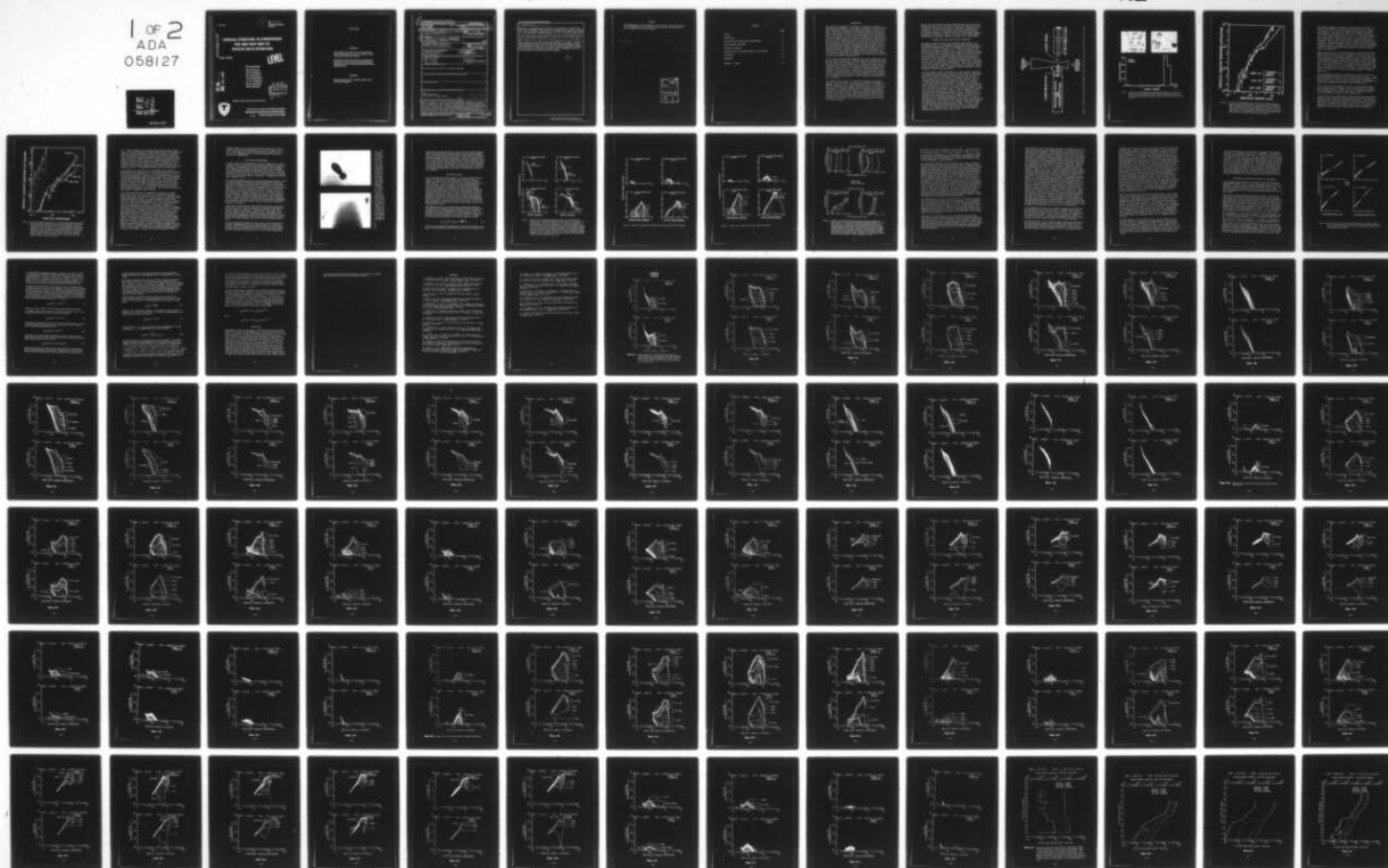
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VERTICAL STRUCTURE IN ATMOSPHERIC FOG AND HAZE AND ITS EFFECTS ON IR EXTINCTION

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Atmospheric Sciences Laboratory

White Sands Missile Range, N.M. 88002

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| 20. ABSTRACT (Continue on reverse side if necessary and identify by block number) Vertical structure of the size distribution and number concentration of particulates in atmospheric fog and haze near Grafenwöhr, West Germany, was measured with a balloon-borne light-scattering aerosol counter for periods spanning parts of 8 days in February 1976. The work was done as part of the Atmospheric Sciences Laboratory's contribution to a US Army field exercise conducted by the Night Vision Laboratory for the purpose of determining the effect of atmospheric constituents on various night vision systems. For haze (5 km visibility) conditions, little vertical variation is seen; but for low visibility (< 1 km) fog, | | |

20. ABSTRACT (cont)

micrometers

conditions, significant vertical increases in concentration of droplets with radii larger than $4\mu\text{m}$ are seen over the first 150 m altitude. For haze, the particle size distribution is approximated by a lognormal with geometric mean radius $r_g \approx 0.2\mu\text{m}$ and geometric standard deviation $\sigma_g \approx 1.9$. For fog, a bimodal distribution is found with a relative maximum for the larger particle mode at radii of $4\mu\text{m}$ to $6\mu\text{m}$ and corresponding values $r_g \approx 5\mu\text{m}$ and $\sigma_g \approx 1.6$; the smaller particle mode has values of $r_g \approx 0.3\mu\text{m}$ to $r_g \approx 0.6\mu\text{m}$ and $\sigma_g \approx 1.8$ to $\sigma_g \approx 2.5$. Liquid water content values for haze and fog range from 10^{-4} to 0.45 g/m^3 . Extinction calculated from the particle size distributions shows an approximate $1/\lambda$ wavelength dependence for haze conditions, but nearly neutral (wavelength independent) extinction for heavy fog. A correlation exists between calculated particulate extinction and calculated liquid water content, independent of particle size distribution, for the fogs and hazes studied.

*0.0001**cu/m*
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PREFACE

We acknowledge Mr. Lucien Biberman of the Institute for Defense Analysis for his suggestion that we look into the correlation of extinction in the infrared and liquid water content.

→ *Lamba*

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CONTENTS

| | <u>Page</u> |
|---|-------------|
| PREFACE | 1 |
| INTRODUCTION | 3 |
| AEROSOL DETECTOR DESCRIPTION AND CALIBRATION | 4 |
| DATA COLLECTION PROCEDURES | 11 |
| HAZE AND FOG RESULTS | 13 |
| EXTINCTION AND LIQUID WATER CONTENT OF FOG AND HAZE | 21 |
| CONCLUSIONS | 25 |
| REFERENCES | 27 |
| APPENDIX. FIGURES | A-1 |

INTRODUCTION

The presence of atmospheric fog and haze is important to radiative transfer in the atmosphere. For example, fog and haze particles degrade the performance of electro-optical systems and affect the earth's climate. Quantitative estimates of these effects generally require knowledge of the vertical variation of fog and haze particle size distributions. Near-surface size distribution measurements have been made by impaction techniques [1-6] by collection of particles on spiderwebs [7], by a laser hologram camera [8], and by light-scattering single-particle counters [9,10]. However, measurements of the vertical structure of fog and haze particle size distributions are difficult to make and consequently are rather sparse with some exceptions [11,12]. Both Pilié et al. and Goodman used impaction techniques to measure droplet spectra to 150-200 m altitudes. Impaction measurements involve laborious microscope measurement of droplets or droplet replicas which restrict their usefulness, as only a relatively small number of distributions can realistically be determined. Furthermore, corrections must be applied to obtain true droplet size and collection efficiencies must be known to obtain relative particle concentrations.

In this report we present measurements of the size distributions of fog and haze particles made with a tethered balloon-borne light-scattering particle counter, with emphasis on their effect on radiation for visible through middle infrared wavelengths. The primary advantage of using a light-scattering counter over an impactor technique is that the measurement is done in situ in real time, with minimal disturbance to the particles during measurement. Measurement of vertical structure of atmospheric fog and haze under relatively stable conditions is possible with this technique because a statistically significant size distribution measurement can be completed in approximately 1/2 minute, permitting an upleg and downleg profile to 200 m altitude to be completed in 15 to 30 minutes.

The purpose of this report is to present the vertical structure of fog and haze that occurred in a wintertime continental environment in Germany, as measured with a light-scattering counter, in the form of particle number, area, and volume distributions, and in the form of extinction and liquid water content values calculated from the measured distributions. An additional purpose is to investigate the spectral dependence of extinction, and to examine correlations between extinction and liquid water content. Measurements reported here were made during February 1976 near Grafenwöhr, a town located 100 km north of Nürnberg near the eastern border of West Germany. The local terrain consists of rolling hills and is partially forested; some of the land is tilled for farming.

Because the light-scattering counter technique offers only an indirect measure of particle size, and because the particular instrument we used is heretofore untested, we first give in section 2 a discussion of its calibration. The balloon experiment data collection procedures are described in section 3. In section 4 typical haze and fog profiles are presented, together with discussion of their significance regarding propagation of radiation. Finally, in section 5 the relation between extinction and liquid water content of fog and haze is investigated.

AEROSOL DETECTOR DESCRIPTION AND CALIBRATION

The aerosol detector used for the measurements reported here is the Knollenberg (after R. G. Knollenberg, the developer) Classical Scattering Aerosol Spectrometer (CSAS) manufactured by Particle Measurement Systems, Inc. (PMS), of Boulder, CO. A schematic of the optical system of this instrument is shown in Fig. 1. The device works on the principle that as aerosol particles flow through an illuminated volume, light scattered into a particular (near forward scattering) solid angle by a single particle is measured photoelectrically and is used to determine particle size by electronically classifying response pulses according to their magnitude. The response signal can be related to particle size by a calibration supplied by the manufacturer. One problem with using the manufacturer calibration is that it does not consider the refractive index of the particle measured. Particles of the same size but of different composition give significantly different response pulses; therefore, we deem it necessary to address the question of the instrument calibration in some detail.

In order to make a definitive measure of the CSAS response characteristics, monodisperse aerosols with known sizes and indexes of refraction are needed. We generated uniform spherical particles of nigrosin dye (index = $1.67-0.26i$, $\lambda = 6328 \text{ \AA}$) with a technique described earlier [13] and measured particle sizes using electron microscopy. We also generated uniform particles of polystyrene latex (index = $1.592-0i$, $\lambda = 5893 \text{ \AA}$) available from Dow Chemical, by atomizing the hydrosol samples (diluted with distilled water). Measurement of uniform aerosol with the CSAS results in a peak in the pulse height spectrum; the position of the peak is a measure of the response of the instrument to aerosol of that size and refractive index. A typical spectrum for monodisperse aerosol of nigrosin dye is shown in Fig. 2, together with a scanning electron microscope micrograph of several of these particles collected onto a Nuclepore filter. The response of the CSAS to solid particles of nigrosin dye and polystyrene latex are shown in Fig. 3. Typical errors in response measurements are also shown. The errors in measurement of particle radii are less than the size of the dots marking the measurements. The squares in Fig. 3 denote response measurements for nearly monodisperse crown glass beads (index = $1.51-0i$, $\lambda = 5893 \text{ \AA}$) available from Particle Information Services, Inc., Grants Pass, Oregon. The standard deviation in particle size of these beads is shown by the horizontal "error" bars, and the standard deviation in response pulses by the vertical "error" bars.

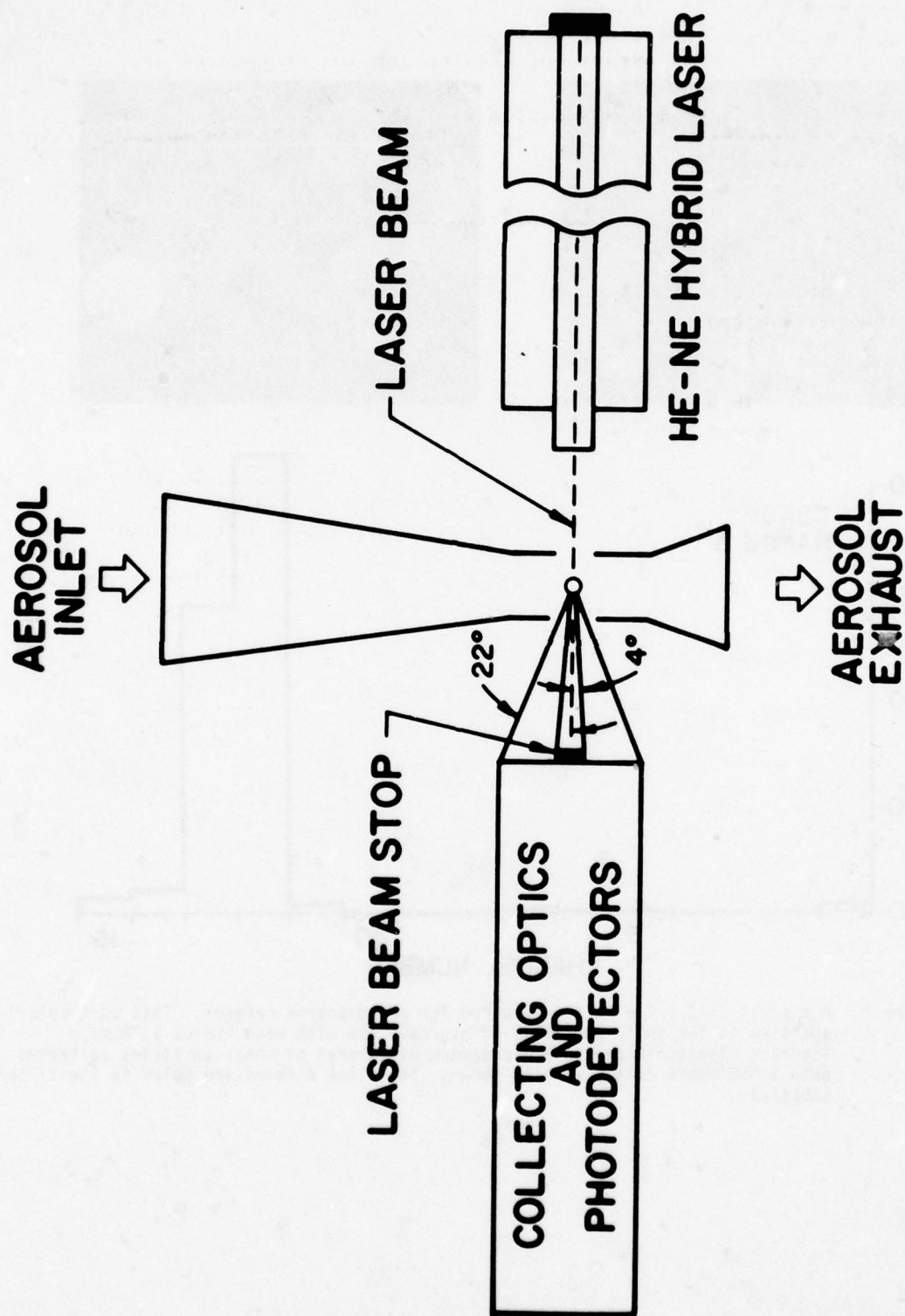


Figure 1. Schematic of the Particle Measurement Systems CSAS light-scattering aerosol counter optical system.

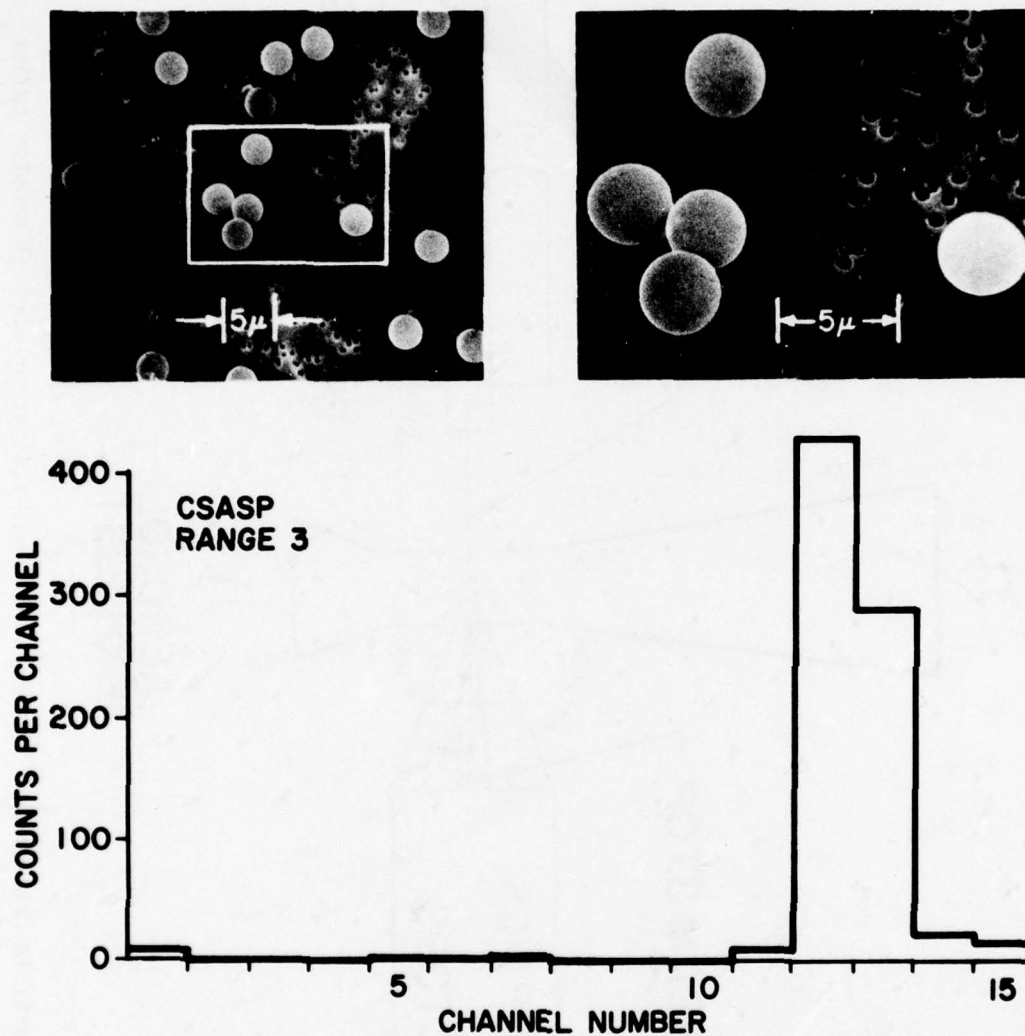


Figure 2. A typical CSAS pulse height spectrum for monodisperse aerosol. This particular spectrum is for solid particles of nigrosin dye with mean radius $1.78\mu\text{m}$; a scanning electron microscope micrograph of several of these particles collected onto a Nuclepore filter is also shown. The black circles are holes in the filter substrate.

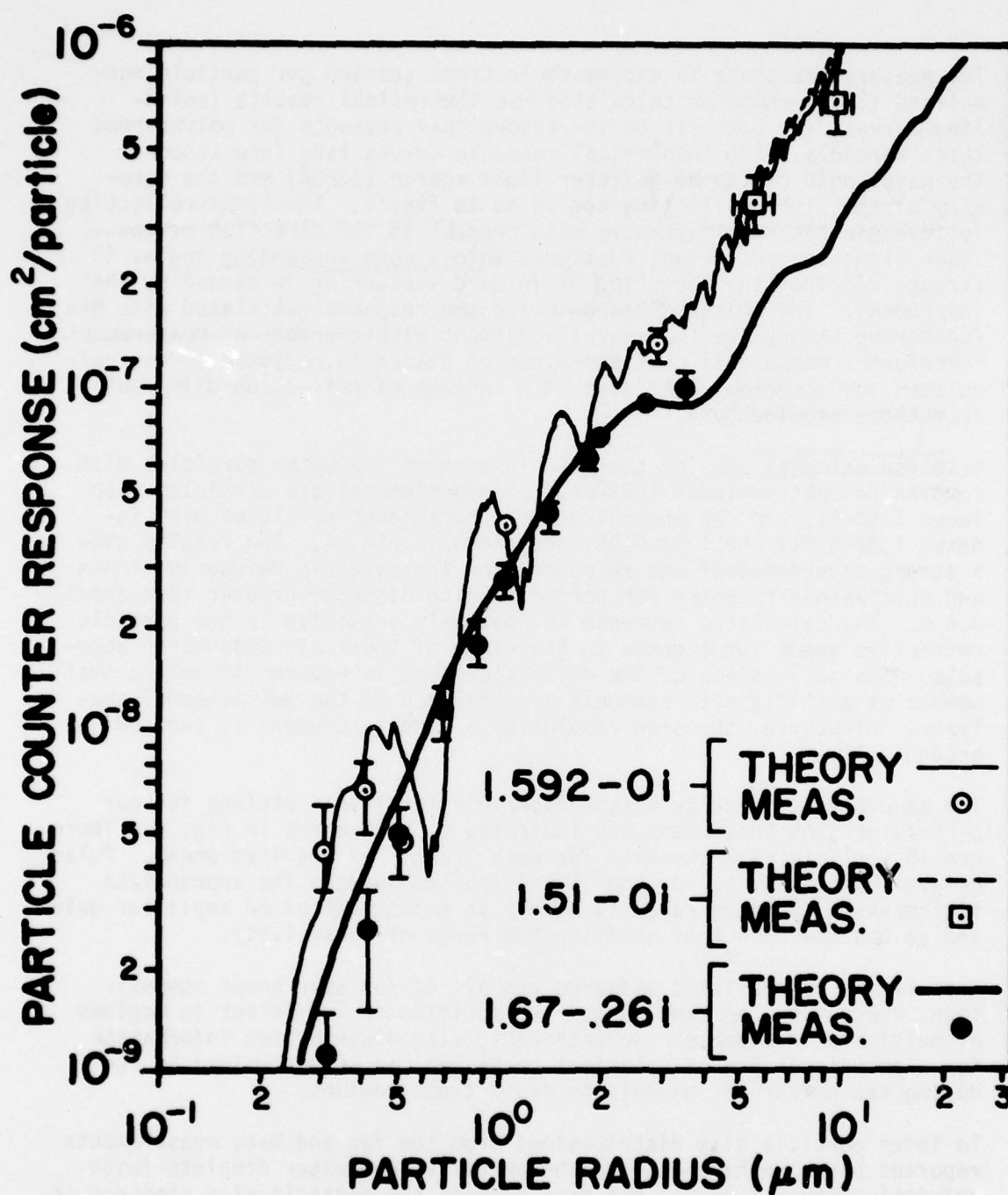


Figure 3. The Particle Measurement Systems CSAS particle counter response; measured (circles and squares) and calculated using Mie scattering theory (smooth curves) for single spherical particles vs particle size. Measured values have been normalized for the best fit to the theoretical values for polystyrene latex particles with refractive index 1.592-0i. The theoretical curve for glass beads with refractive index 1.51-0i extends only to about 4μm radius.

The measured response is expressed in cross section per particle normalized to the computer calculated Mie theoretical results (solid-line curves) for best fit to the theoretical response for polystyrene latex aerosols. The theoretical response curves take into account the wavelength of the He-Ne laser light source (6328Å) and the geometry of the light-collecting optics as in Fig. 1. The light-collecting solid angle has axial symmetry with respect to the direction of the laser light source. Light scattered into a cone subtending angles 4° through 22° from the direction of forward scattering is sensed by the instrument. The measured response and the response calculated with Mie scattering theory are in general agreement within errors of measurement. Therefore a measure of confidence can be placed in response curves calculated for spherical particles with indexes of refraction different from those studied here.

Response calculations for the CSAS instrument for water particles with complex refractive index $1.33-0i$, for ammonium sulfate particles with index $1.50-0i$, and for absorbing atmospheric dust particles with indexes $1.50-0.01i$ and $1.50-0.05i$ are shown in Fig. 4. The results show a strong dependence of the response upon the particle refractive index and multivalued response for particles with diameter greater than about $0.8\mu\text{m}$. The calculated response is obviously sensitive to the particle refractive index for indexes in the range of those of atmospheric aerosols. The seriousness of the multivaluedness is reduced if only a small number of particle size channels are utilized in the multichannel analyzer. Of course, the size resolution of the instrument is reduced accordingly.

The manufacturer's pulse height discriminator level settings for our particular CSAS instrument are indicated by tick marks in Fig. 4. There are 15 particle size channels for each "range" of the instrument. Pulse height channels 1, 5, 10, and 15 are labeled between the appropriate tick marks. Changing range is merely an adjustment of an amplifier gain and so has the effect of shifting the range of sensitivity.

Even if the particulates measured are all of the same known composition, there are - on some ranges - discriminator levels set in regions of multivalued response. Nevertheless, size distribution information for a polydispersion of spherical particles can be determined by reducing the number of channels to avoid these regions.

To infer particle size distributions from the fog and haze measurements reported here, we have assumed the particles are water droplets (with refractive index $1.33-0i$) and have grouped the particle size channels to avoid regions of multivalued response. This channel grouping for water particles is indicated by the heavy tick marks in Fig. 4. Specific channel size limits for the different ranges of the instrument can be determined from this figure by noting at what radii on the water

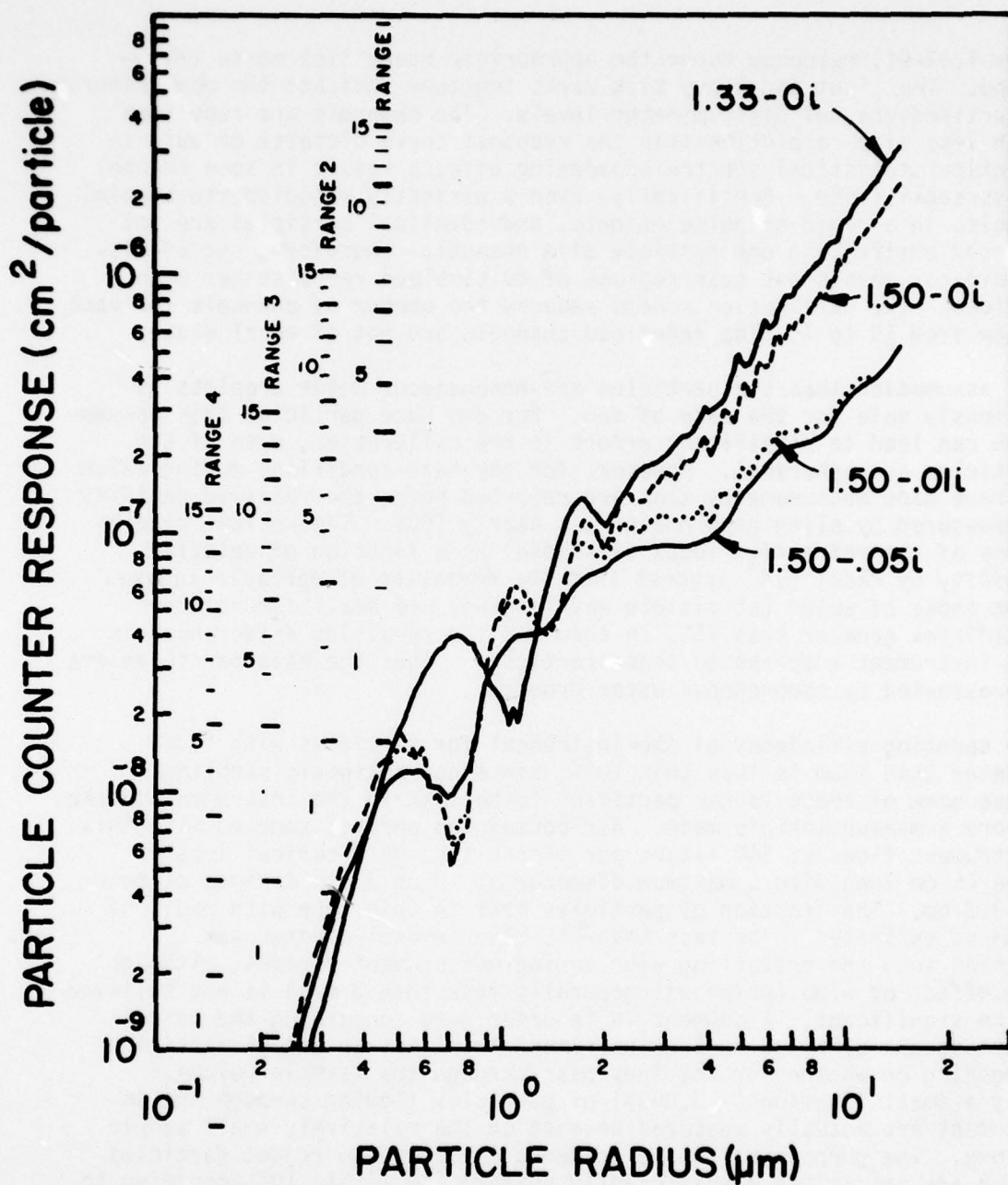


Figure 4. Mie theory response calculations for the Particle Measurement Systems CSAS particle counter for water particles with refractive index 1.33-0i, ammonium sulfate with approximate index 1.50-0i, and atmospheric dust with indexes 1.5-0.01i and 1.50-.05i. The tick marks indicate the pulse height discriminator level settings as set by the manufacturer for the detector. Channels 1, 5, and 15 are labeled between the appropriate tick marks for the different range settings of the instrument. The heavy tick marks indicate the pulse height discriminator level settings used in grouping channels together to avoid regions of multivalued response under the assumption that all particles are water.

($m = 1.33 - 0i$) response curve the appropriate heavy tick marks correspond. The light and heavy tick marks together indicate the manufacturer advertised channel discriminator levels. The channels are redefined with less size resolution than the response curve dictates because in practice statistical spectra broadening effects result in some channel cross sensitivity. Specifically, even a perfectly monodisperse aerosol results in a range of pulse heights, and identical particles are not counted entirely in one particle size channel. Therefore, use of discriminator levels set near regions of multivalued response has been avoided. The calibration scheme reduces the number of channels for each range from 15 to 7. The redefined channels are not of equal width.

Our assumption that the particles are homogeneous water droplets is obviously safe for the case of fog. For dry haze particles this assumption can lead to significant errors in the calibration, even if the particles are spherical. However, for the haze conditions during which we have made measurements that are reported here, the relative humidity as measured by sling psychrometer was nearly 100%. Theoretical predictions of the refractive index of aerosol as a function of relative humidity by Hänel [14] suggest that the deviation of particle indexes from those of water (at visible wavelengths) are small for relative humidities greater than 95%, in terms of the resulting differences in the instrument response to these particles. Thus the haze particles are approximated by homogeneous water droplets.

The counting efficiency of the instrument for particles with radii greater than $15\mu\text{m}$ is less than 100%, since nonisokinetic samplings cause some of these larger particles to be lost in the instrument intake before a measurement is made. Air containing aerosol sampled with this instrument flows at 340 liters per minute through a conical intake tube 45 cm long with a maximum diameter of 10 cm and a minimum diameter of 3.3 cm. The fraction of particles lost in this tube with radii of $15\mu\text{m}$ is estimated to be less than 7%. The aerosol counter was pointed into the prevailing wind during measurement periods, although the effect of wind (which was generally less than 3 m/s) is not believed to be significant. A comment is in order here concerning the coincidence scheme utilized in the instrument to reject or accept particles depending on whether or not they pass through the "sample volume." Only a small fraction ($\sim 0.003\%$) of particles flowing through the instrument are actually measured because of the relatively small sample volume. The purpose of the coincidence scheme is to reject particles which are not within a sufficiently uniform (to within 10% according to PMS) part of the laser beam by opto-electronically discriminating against out-of-focus particles. There is evidence to suggest that this scheme results in a sample volume that is somewhat dependent on particle sizes. In other words, the instrument flow rate may be different for different size channels. However, simultaneous measurements made on

uniform aerosols in our laboratory with both the CSAS and a particle counter of a special design developed by Rosen [15] suggest this effect is less than 20%, at least for particles with radii less than $1\mu\text{m}$. A more complete account of the aerosol detector calibration problem is in preparation.

DATA COLLECTION PROCEDURES

The basic goal of the experiment was to obtain information on vertical profiles of particle size distribution during atmospheric fog and haze conditions. To accomplish this, the CSAS particle counter together with temperature and visible radiation flux sensors was incorporated into a single payload weighing approximately 9 kg and carried aloft to a maximum altitude of 250 m by a 1500 ft³ tethered balloon. Digitized data was sent via a wire link to a data acquisition unit on the ground; this link also furnished power to the payload. A photograph of the system is shown in Fig. 5.

This experiment was done as part of a much larger field exercise, and as a result considerable constraint on operating schedules was imposed. For example, there was a problem caused by presence of a nearby radar station, so that data could only be collected when the radar was not in operation. In addition, the balloon experiment could not be operated at night because it was not equipped with lights which were required to meet local aircraft hazard warning regulations. Choice of actual operating conditions (ascent and descent rates, and altitude intervals) was determined by a compromise between the need for long counting intervals to produce statistically meaningful aerosol size distribution results, the desire for short counting intervals to obtain reasonable vertical spacial resolution, and the need to complete the ascent/descent in a relatively short time interval because of continually changing atmospheric conditions.

A choice was also made in the field controlling the CSAS counter size range (see Fig. 4). Under the assumption of spherical water droplets the size ranges are: range 1, 0.5-16 μm radius; range 2, 0.37-8 μm radius; range 3, 0.29-4 μm radius; range 4, 0.24-1.6 μm radius. Since the range setting could not be changed remotely, measurements made during an upleg and downleg traverse were necessarily made for a single range setting, optimized to cover the most important range of particle size for the widely different fog and haze size distributions encountered.

The usual operational procedure for a particular balloon traverse was to sample intermittently (with the CSAS positioned at a fixed altitude and set on 30- to 60-second accumulation times) at 30 m altitude intervals during the upleg portion, and to sample continuously (with the CSAS continuously moving and set on 20- to 30-second accumulation

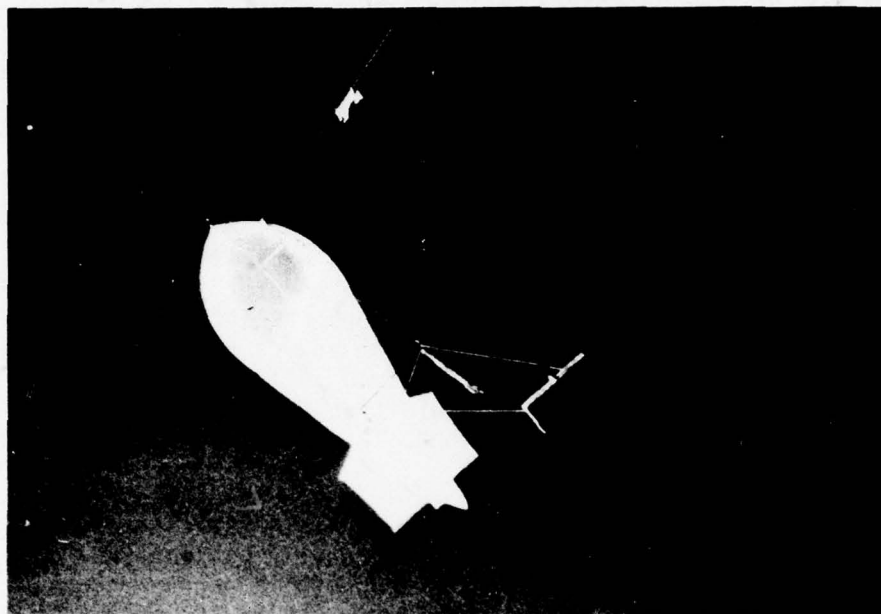
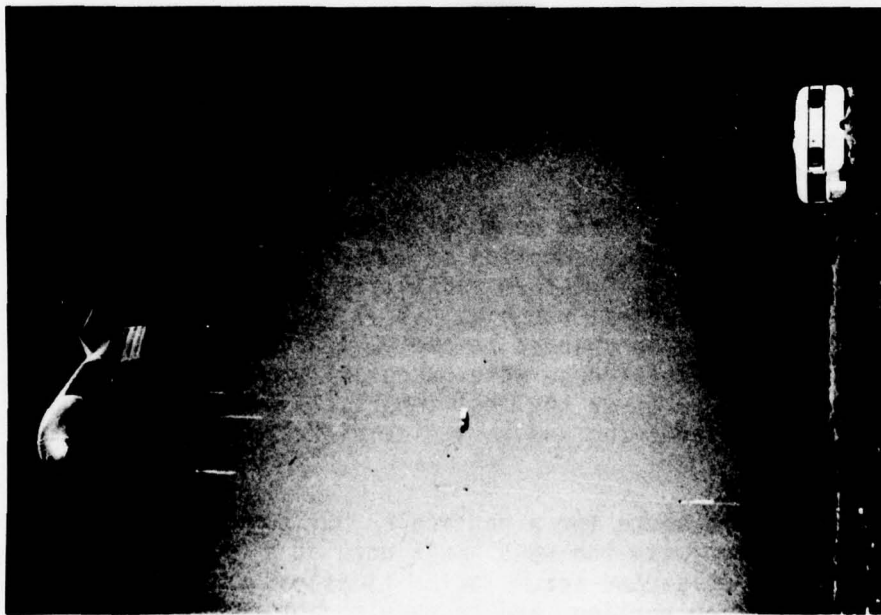


Figure 5. Photographs of the balloon-borne aerosol detector system used for fog and haze measurements near Grafenwöhr, West Germany. The detector (a light-scattering aerosol counter model CSAS manufactured by Particle Measurement Systems of Boulder, CO) is suspended approximately 15 m below the 1500 ft³ tethered balloon and is rigged so that the detector intake points into the prevailing wind. Temperature and radiation flux sensors are incorporated into the payload.

times) during the downleg portion. Since the size distribution measurements made during upleg and downleg portions of each traverse compared well, and since more vertical resolution was obtained during the downleg portion, we concentrate only on the latter. Thus, particle size distribution measurements reported here have been obtained by integrating over 8 m to 12 m altitude intervals (depending on the CSAS accumulation time), as the payload descent rate was about 0.4 m/s. The altitude identified with a particular size distribution is the lower limit of the corresponding altitude interval. During heavy fog conditions it was not possible to achieve altitudes greater than 150 m, as copious amounts of water and rime condensed onto the balloon and tether line, reducing the balloon lift capability.

HAZE AND FOG RESULTS

Four profiles of haze and fog, typical of 19 profiles measured during 8 days in February 1976 near Grafenwöhr are presented in Figures 6 through 8. These show the vertical structure in terms of particle number ($dN/d \log r$), cross sectional ($dS/d \log r$), and volume ($dV/d \log r$) distributions. Some of the particle counter measurements (solid lines) have been fitted with lognormal size distributions (dashed curves) for the purpose of estimating the effects of extending and smoothing the measured size distributions. The reason for showing the area and volume distributions is that for limiting regimes of particle size the extinction cross section is nearly proportional to either the integrated cross-sectional area (for particles large compared to the wavelength) or volume (for particles small compared to the wavelength, as will be discussed in a following section). These distributions can therefore be used to infer the range of particle sizes that contribute most to extinction. Generally the values for $dN/d \log r < 10 \text{ cm}^{-3}$ are based upon fewer than 10 particle counts per size channel and thus have large statistical errors.

Corresponding profiles of the liquid water content (LWC) and extinction coefficients calculated from the measured and fitted distributions at $0.55 \mu\text{m}$, $4 \mu\text{m}$, and $10 \mu\text{m}$ wavelengths are shown in Fig. 9. Note that the abscissa scales are different for the different parts of Fig. 9. The LWC and extinction for the fitted size distributions are calculated only for one or two altitudes for each profile in Fig. 9. The Mie single scatter extinction coefficients were calculated as follows:

$$\beta_{\text{ext}} = \int_{-0}^{\infty} \pi r^2 Q_{\text{ext}}(m, x) \frac{dN(r)}{dr} dr \quad (1)$$

Here $N(r)$ is the particle concentration for particles with radius $\geq r$ and Q_{ext} is the Mie efficiency factor defined as the ratio of single particle

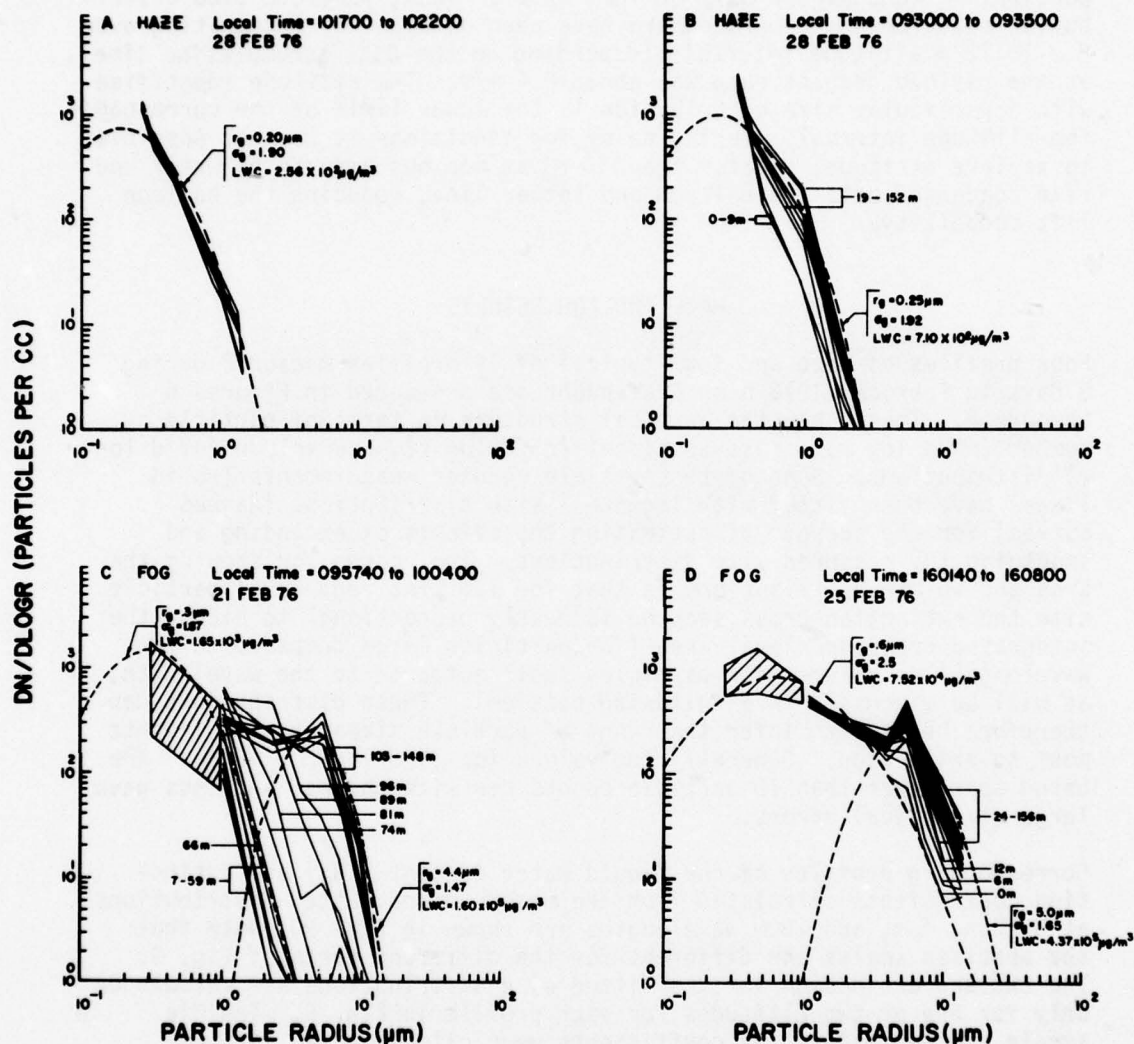


Figure 6. Particulate size distribution measurements made near Grafenwöhr, West Germany, with a tethered balloon-borne aerosol counter for several haze and fog conditions. Each size distribution is for a particular altitude or altitude range as indicated. Measurements at particular altitudes (121 m in Fig. 6A, 95 m in Fig. 6B, 59 m and 126 m in Fig. 6C, and ground level and 135 m in Fig. 6D) have been fitted with lognormal size distributions (dashed curves) for the purpose of estimating the effects of extrapolating the measured distributions. Values of geometric mean radius r_g , geometric standard deviation σ_g , and liquid water content LWC, are shown with the corresponding lognormal curves. The date, local time (hours, minutes, seconds), and time interval for the measurements are given.

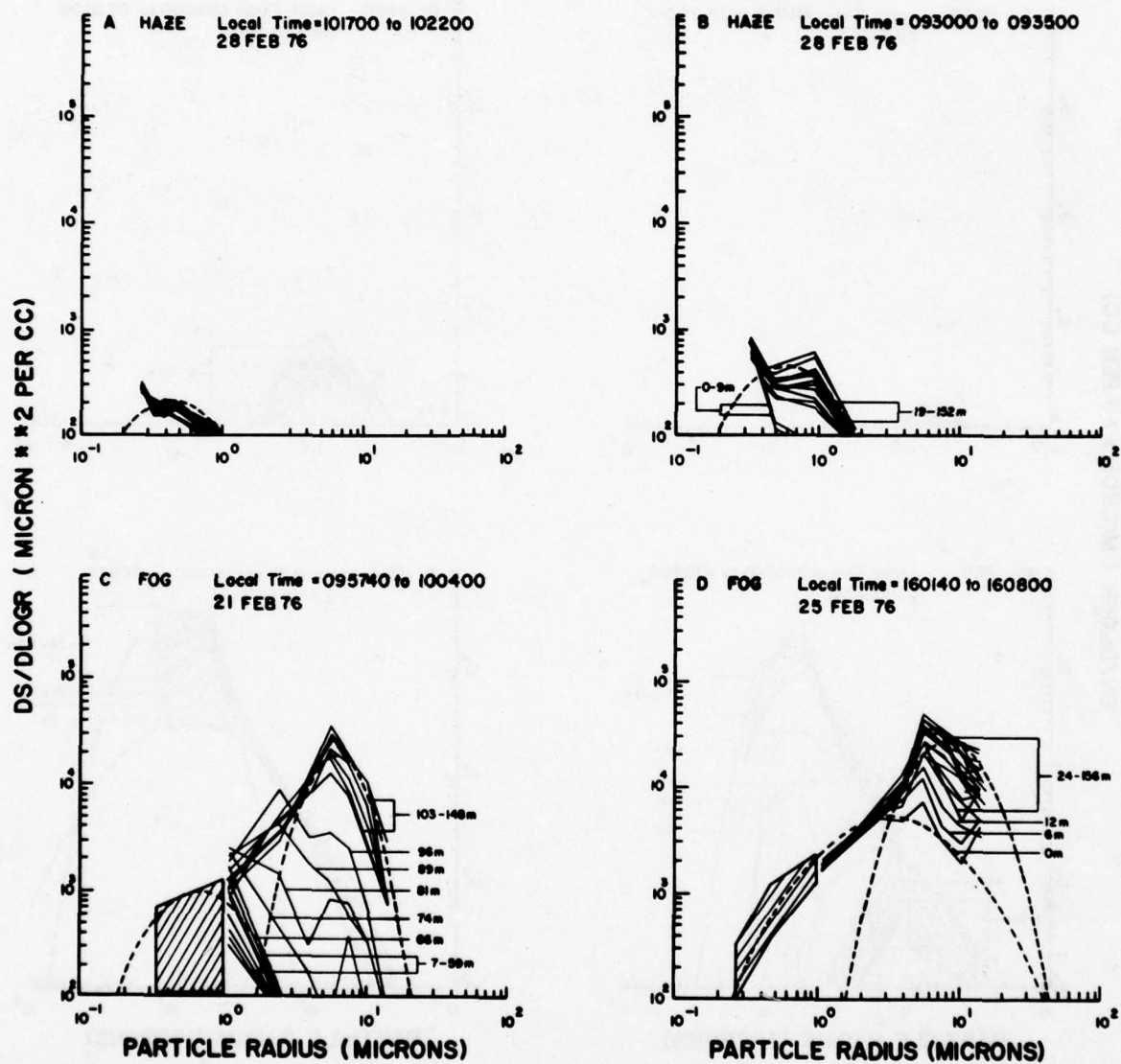


Figure 7. Same as Fig. 6 except for particle area (cross-sectional) distribution.

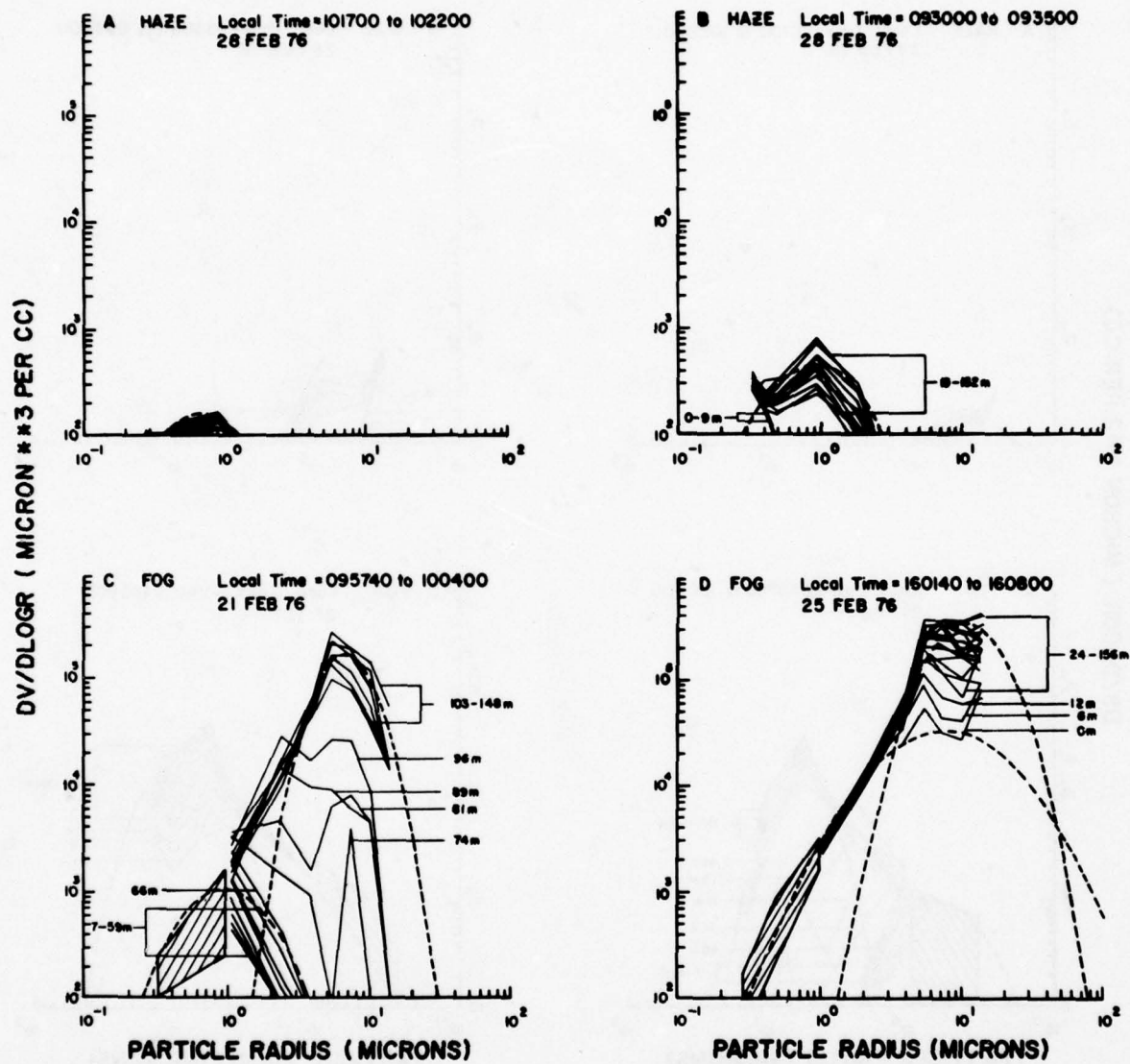


Figure 8. Same as Fig. 6 except for particle volume distribution.

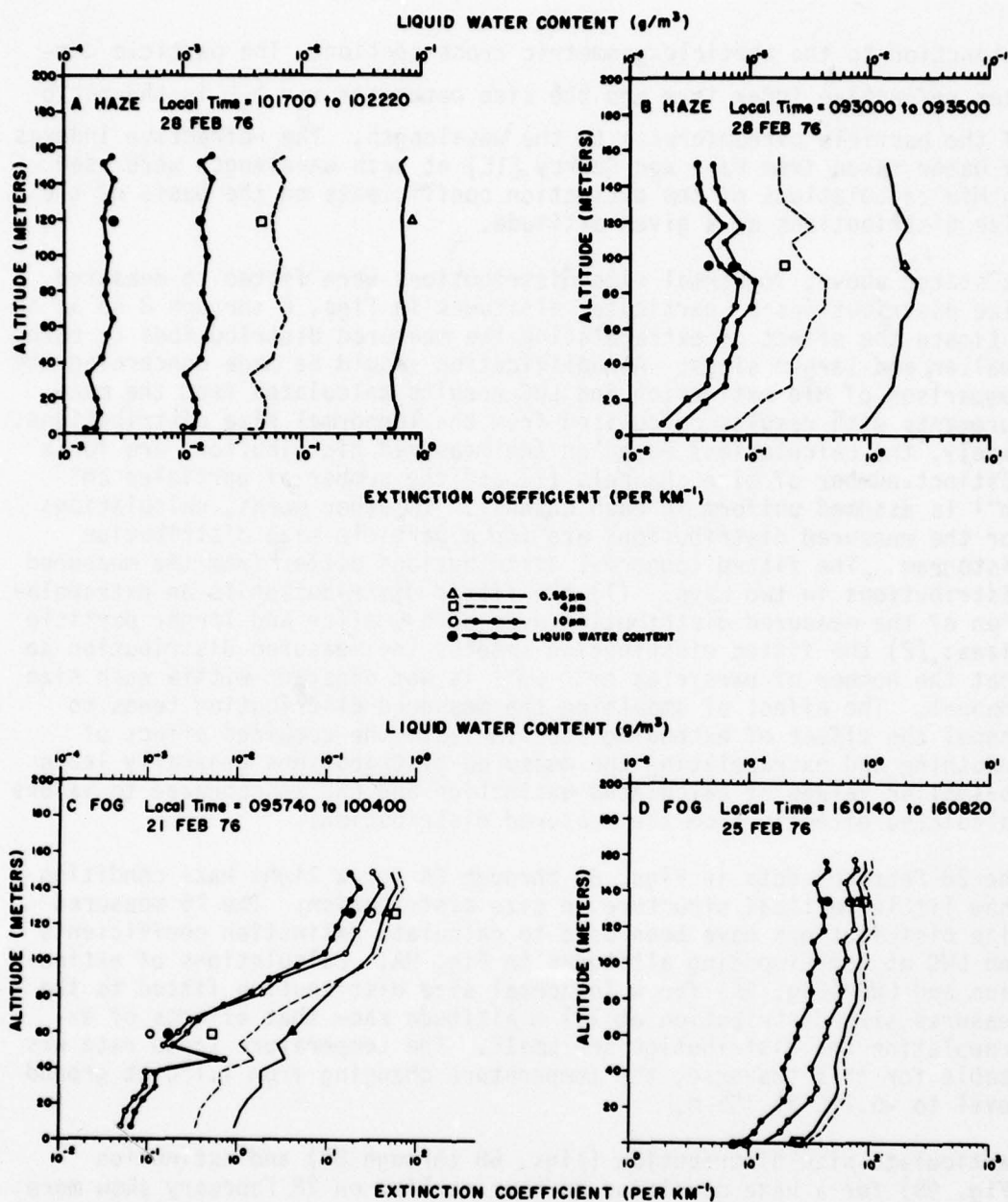


Figure 9. Profiles of particulate extinction and water content calculated from particle size distribution and concentration measurements shown in Fig. 6. The solid-dot curve indicates the liquid water content and the solid, dashed, and open-dot curves indicate the extinction at 0.55 μm , 4 μm , and 10 μm wavelengths. Calculations of the extinction and LWC for particular lognormal distributions fitted to measured distributions (Figs. 6 through 8) are shown at the corresponding altitudes, and are denoted by triangle, square, open-circle, and solid circle symbols for extinction at 0.55 μm , 4 μm , 10 μm , and liquid water content. The date, local time (hours, minutes, seconds), and time interval for the measurements are given.

extinction to the particle geometric cross section. The particle complex refractive index is m and the size parameter $x = \frac{2\pi r}{\lambda}$ is the ratio of the particle circumference to the wavelength. The refractive indexes of water taken from Hale and Querry [16] at each wavelength were used in Mie calculations of the extinction coefficients on the basis of the size distributions at a given altitude.

As stated above, lognormal size distributions were fitted to measured size distributions at particular altitudes in Figs. 6 through 8 so as to estimate the effect of extrapolating the measured distributions to both smaller and larger sizes. A qualification should be made concerning the comparison of Mie extinction and LWC results calculated from the measurements with results calculated from the lognormal size distributions. Namely, the calculations based on the measured distributions are for a distinct number of size channels (7) and the number of particles $\text{cm}^{-3} \mu\text{m}^{-1}$ is assumed uniform in each channel. In other words, calculations for the measured distributions are for a particle size distribution histogram. The fitted lognormal distributions differ from the measured distributions in two ways: (1) the fitted distribution is an extrapolation of the measured distribution over both smaller and larger particle sizes; (2) the fitted distribution smooths the measured distribution so that the number of particles $\text{cm}^{-3} \mu\text{m}^{-1}$ is not constant within each size channel. The effect of smoothing the measured distribution tends to cancel the effect of extending it. In fact, the combined effect of smoothing and extrapolating the measured distributions generally leads to smaller values of calculated extinction and LWC as compared to values calculated directly from the measured distribution.

The 28 February data in Figs. 6A through 8A for a light haze condition show little vertical structure in size distribution. The 16 measured size distributions have been used to calculate extinction coefficients and LWC at corresponding altitudes in Fig. 9A. Calculations of extinction and LWC (Fig. 9A) for a lognormal size distribution fitted to the measured size distribution at 121 m altitude show that effects of extrapolating the distribution are small. The temperature lapse rate was stable for this traverse, the temperature changing from 1.1°C at ground level to -0.1°C at 155 m.

Particulate size distribution (Figs. 6B through 8B) and extinction (Fig. 9B) for a haze condition an hour earlier on 28 February show more vertical variation. A visual estimate of 2 km horizontal visibility was made during this traverse. Using Koschmieder's formula to relate visibility to extinction for a threshold contrast ratio of 2%, this yields a value of 1.96 km^{-1} extinction for this haze condition, comparing favorably with extinction at $0.55 \mu\text{m}$ (Fig. 9B) calculated from the measured size distributions.

The fog data in Fig. 6C show large increases in concentration of $4\mu\text{m}$ to $16\mu\text{m}$ radius particles with altitude and Figs. 7C, 8C, and 9C show the effects of these increases. The cross-hatched areas indicate the altitude variation in measurements made during traverses before and after the traverse in question, with the instrument set so as to sense smaller particles. Assuming that temporal changes are relatively small during these series of traverses (a period of about 2 hours for all traverses was involved here), the cross-hatched areas show the approximate shape of the envelope of the distribution for smaller particles. Generally, the concentration of these smaller particles increased with altitude. The temperature lapse rate for this traverse is again stable with a ground temperature of -3.3°C . The calculated extinction coefficients and LWC in Fig. 9C vary by 2 to 3 orders of magnitude over this 150-m traverse. Comparison of results from the measured and fitted lognormal size distributions suggests that effects of smoothing and extrapolating the measured distribution are significant (as much as a factor 2 in extinction at $10\mu\text{m}$, but nearly zero at $0.55\mu\text{m}$) for altitudes less than about 70 m. At higher altitudes, the effects of smoothing and extrapolating the size distributions would appear to be relatively small, since at 126 m altitude the differences in extinction and LWC are less than 5%. For the largest particle size channel (centered at $14\mu\text{m}$ radius) the number of counts for this traverse is not statistically significant. In fact, the point to which six number, area, and volume size distribution curves (at altitudes 111-148 m) converge in Figs. 6C through 8C correspond to only 1 particle count for each curve. Nevertheless, the trend in the size distributions for the next two smaller size channels (centered at $12\mu\text{m}$ and $9\mu\text{m}$) strongly suggests the measured fog distributions can be adequately extrapolated to larger particle sizes with the lognormal. Further, the shape of the distribution for the larger particle sizes is in good agreement with the preceding up-leg measurements made about 15 minutes earlier. We point out here that the measured distribution at 126 m altitude was actually fitted by a sum of three lognormal distributions (two of which are shown here) to better accommodate the measurements in the $1\mu\text{m}$ to $4\mu\text{m}$ radius interval.

It is noteworthy that during this traverse a visual siting of the balloon payload was lost at 148 m altitude. Integrating the $0.55\mu\text{m}$ extinction coefficient profile over the first 148 m of this traverse, we find an optical depth of 3.02, a value 23% lower than expected from the Koschmieder relation. Considering the uncertainty in making a visibility estimate and in applying the Koschmieder relation to relate visibility to optical depth, we judge this to be good agreement.

The heavy fog data in Fig. 9D shows increasing extinction coefficient with altitude, again caused by the increased concentration of $4\mu\text{m}$ to $16\mu\text{m}$ radius particles shown in Figs. 6D through 8D. The 21 size distribution curves in Figs. 6D through 8D were used to calculate extinction coefficients and LWC presented in Fig. 9D. In contrast to the

21 February fog, vertical increases in extinction and LWC are only about a factor 2 to 4. Again the cross-hatched areas in Figs. 6D through 8D show the shape of distributions for submicrometer particles for a balloon traverse made 1/2 hour after the traverse in question. This traverse is similar to six traverses made during stable fog conditions 25-26 February 1976 in Grafenwöhr. Ground windspeed for each of these traverses was generally less than 2 m/s, suggesting the fog was radiation type rather than advection type. Here it is evident that the log-normal extrapolation to particle sizes larger than those measured may be significantly in error, particularly in calculations of LWC (Fig. 8D) and extinction at $10\mu\text{m}$, for which a third mode of larger particles may contribute significantly, even though their relative number may be small. However, although a third mode of larger fog particles cannot be ruled out, a bimodal nature of the distributions is in agreement with measurements of wintertime radiation fogs in England by Garland et al. [1,17] by Roach et al. [18] and some radiation fog measurements by Krasikov and Chikirova [2] near Leningrad. The extinction coefficient is neutral (wavelength independent) within a factor of about 2; the extinction at $4\mu\text{m}$ being slightly greater than that at $.55\mu\text{m}$ and $10\mu\text{m}$. Errors due to size distribution extrapolation and smoothing, assuming no mode of particles with radii greater than $16\mu\text{m}$ is important, are small. The nearly neutral extinction for low visibility conditions has been found by Patterson [19] for soil-derived aerosols as well, suggesting that a broad range of low (< 1 km) visibility atmospheric conditions result in neutral extinction from visible wavelengths through the middle-infrared ($0.5\mu\text{m}$ to $10\mu\text{m}$).

Each of the four haze and fog profiles presented here is for a downleg portion of a balloon traverse. The corresponding upleg values of extinction and liquid water content are generally within 50% of the downleg values, so that temporal changes encountered are usually small compared to the vertical variations. These four profiles are typical of the 19 profiles taken during 8 days at Grafenwöhr. For haze conditions, little vertical variation in the size distribution is observed; but for fog conditions, there are significant increases in the concentration of droplets with radii greater than about $4\mu\text{m}$. Measurements of the downward visible radiation flux made with a photodetector attached to the balloon package show that for each of the traverses the flux change with altitude was less than 10%. This suggests that the fog or haze depths were somewhat greater than the maximum altitudes attained.

The trend toward increasing concentration of larger droplets with increasing altitude is in agreement with measurements of advection fogs over San Francisco by Goodman [12], although in direct contradiction to previous measurements of continental fogs, notably those of Pilié et al. [11] for fogs in the Chemung River Valley near Elmira, New York. In view of the paucity of definitive measurements of the vertical structure of fog and haze, similar measurements at different geographical locations and under different meteorological conditions are clearly warranted.

For sake of completeness we have given in the appendix the remaining data for all 19 balloon traverses made in Grafenwöhr. Shown are the upleg and downleg values of $dN/d \log r$ vs radius (Fig. A1), $dS/d \log r$ vs radius (Fig. A2), $dV/d \log r$ (Fig. A3), downleg extinction and LWC vs altitude (Fig. A4), and upleg and downleg values of LWC vs altitude (Fig. A5). The data are presented in chronological order; haze conditions existed during 14 December 1975 and 28 February 1976, fog during 21, 22, 23, 25, and 26 February 1976. The volume distributions for the 25-26 February fog data reveal an increasing contribution of larger ($> 10 \mu\text{m}$ radius) particles to LWC and suggest a significant error in LWC resulting from inability of the detector to measure droplets with radius greater than $16 \mu\text{m}$. Temporal variations during these traverses were generally small and less than vertical variations, as is evidenced by comparison of upleg and downleg LWC profiles in Fig. A5.

EXTINCTION AND LIQUID WATER CONTENT OF FOG AND HAZE

To investigate the correlation of particulate extinction and liquid water content evident in Fig. 9, these data, along with data from the remaining 15 balloon traverses made in Grafenwöhr, appear in Fig. 10. Plotted are values of the Mie calculated extinction for $0.55 \mu\text{m}$, $1.2 \mu\text{m}$, $4 \mu\text{m}$, and $10 \mu\text{m}$ wavelengths versus the liquid water content. Each point corresponds to a size distribution measurement made with the aerosol counter.

These results show approximate relationships between extinction and liquid water content, for 289 measured distributions of fog and haze with liquid water content values ranging over four orders of magnitude. Such relationships have been noted before at visible wavelengths by Eldridge [20,21], Barteneva and Polyakova [6], and Kumai [3]. Recently, Chýlek [22] has shown that in fact a linear relationship should exist between extinction and LWC, independent of particle size distribution, but only for a wavelength determined by the radii of the largest particles present in the polydispersion of droplets. This is a result of the efficiency factor for extinction Q_{ext} being nearly proportional to size parameter x , which leads to extinction being proportional to LWC, providing droplets have size parameters less than a maximum value x_m . The maximum size parameter $x_m \approx 7$, but depends slightly on refractive index and hence on wavelength. If we choose a wavelength of $10 \mu\text{m}$, then $x_m = 8$ [22] and according to the Chýlek approximation we should find a linear relation between extinction and LWC for polydispersions of droplets having radii $r \leq 12 \mu\text{m}$. Most of our haze and fog distributions satisfy this condition and, as verified in Fig. 10, there does exist a nearly linear relation between extinction at $10 \mu\text{m}$ and LWC. Even for heavy fog the presence of larger droplets ($r > 12 \mu\text{m}$), insofar as they do not dominate the smaller ones, does not destroy the linear

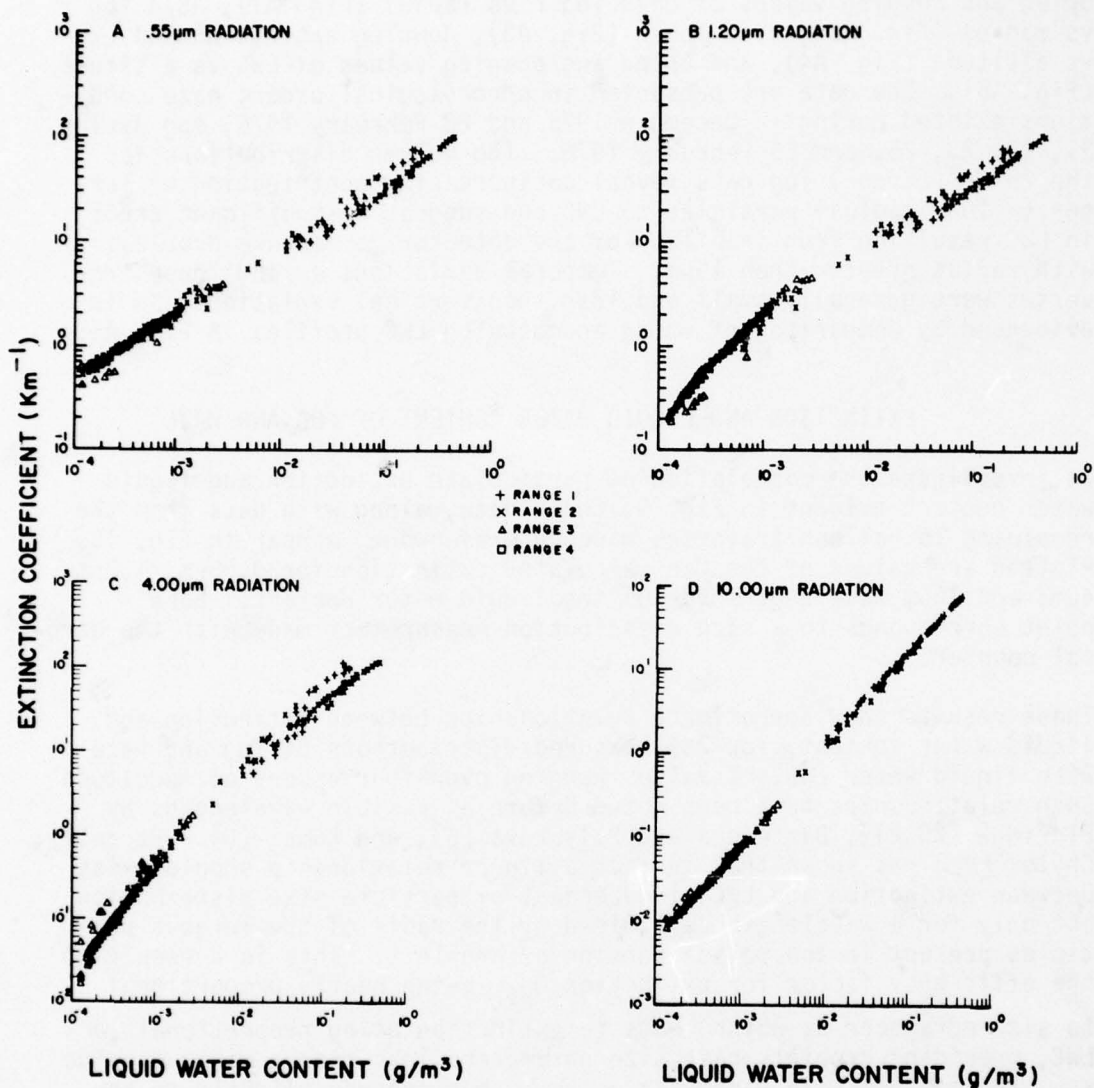


Figure 10. Values of Mie theory calculated particulate extinction at 0.55 μm , 1.2 μm , 4 μm , and 10 μm wavelengths vs liquid water content for a variety of particle size distributions of atmospheric fog and haze measured during February 1976 near Grafenwöhr, West Germany.

size distribution independent relation. However, the fact that there are approximate relationships between extinction and LWC for shorter wavelengths (namely $0.55\mu\text{m}$, $1.2\mu\text{m}$, and $4\mu\text{m}$) for heavy fog (for which particles with $x > 7$ dominate extinction) is probably a result of the common form of the measured fog distributions (see for example, Fig. 6D), since for arbitrary distributions and wavelengths there is no reason to expect a unique relationship between extinction and LWC.

To check our contention that there should not exist a unique relation between extinction and LWC in fog, regardless of the form of the droplet size distribution or value of the wavelength, we compare empirical relationships between extinction at visible wavelengths and LWC determined by other workers for fog measurements made under different meteorological conditions and at different geographical locales. For the present work an approximate empirical relation between extinction at $0.55\mu\text{m}$ and LWC can be derived from the results in Fig. 10A:

$$\beta_{\text{ext}}(0.55) \approx 145 w^{.63}, \quad (2)$$

where $\beta_{\text{ext}}(0.55)$ refers to the particulate extinction coefficient at $0.55\mu\text{m}$, in km^{-1} , and w refers to liquid water content in g/m^3 . For comparison Eldridge [21] gives the empirical relation

$$\beta_{\text{ext}}(0.55) \approx 160 w^{.65}, \quad (3)$$

for fogs and hazes having liquid water content 0.001 - $0.2 \text{ g}/\text{m}^3$. For fogs in the Tsei gorge near Leningrad with liquid water content 0.0002 - $0.2 \text{ g}/\text{m}^3$, Barteneva and Polyakova [6] find

$$\beta_{\text{ext}}(\text{visible}) \approx 142 w^{.87}. \quad (4)$$

From Kumai's [3] simultaneous measurements of arctic fog size distribution and prevailing visual range at Point Barrow, Alaska, with liquid water content 0.005 - $0.1 \text{ g}/\text{m}^3$, we find

$$\beta_{\text{ext}}(\text{visible}) \approx (100 \text{ to } 200) w. \quad (5)$$

where the extinction has been inferred from the visual range measurements using Koschmieder's relation for a threshold ratio of 2%. Kumai has calculated the extinction from the measured size distributions as

we have done and the visual range calculated from these extinction values at $0.55\mu\text{m}$ are in good agreement with his measurements of visual range.

Using these four empirical expressions relating visible extinction to LWC, for an LWC of 0.01 g/m^3 , we get extinction coefficients of 8.0 km^{-1} , 8.0 km^{-1} , 2.6 km^{-1} , and $1.0\text{--}2.0\text{ km}^{-1}$, respectively, for the present work, Eldridge, Barteneva and Polyakova, and Kumai. These values differ by roughly an order of magnitude. These differences are most likely due to the widely differing fog droplet size distributions measured by the various authors, and thus any size distribution independent relation between extinction (at visible wavelengths) and LWC cannot be applied to fogs in general.

The approximation proposed by Chýlek [20] can also be used to explain the approximate $1/\lambda$ wavelength dependence of extinction for haze evident in Figs. 9A and 9B. The approximation can be written in the form

$$\beta_{\text{ext}}(\lambda) \approx \frac{3\pi c}{2\rho\lambda} W \quad (6)$$

where W is the liquid water content, ρ the density of water, and the coefficient c is equal to the slope of a straight line approximating the efficiency factor for extinction by

$$Q_{\text{ext}}(x, \lambda) \approx c(\lambda)x. \quad (7)$$

The coefficient c is a slowly varying function of wavelength. For two wavelengths λ_1, λ_2 the approximation (6) can be rewritten

$$\beta_{\text{ext}}(\lambda_1) \approx \frac{c_1\lambda_2}{c_2\lambda_1} \beta_{\text{ext}}(\lambda_2), \quad (8)$$

relating extinction at one wavelength to that at another. This approximation is only valid for polydispersions of droplets having size parameter less than a maximum value x_m , or equivalently, a maximum radius r_m . From Chýlek's [22] Table 1 the maximum radii and corresponding c values are: at $\lambda = 0.5\mu\text{m}$, $r_m = 0.5\mu\text{m}$, $c = 0.61$; at $\lambda = 3.8\mu\text{m}$, $r_m = 3.6\mu\text{m}$, $c = 0.68$; and at $\lambda = 10\mu\text{m}$, $r_m = 12\mu\text{m}$, $c = 0.35$. Haze particles have maximum radii of about $2\mu\text{m}$ (Figs. 6A and 6B); hence the approximation (8) can be applied at $4\mu\text{m}$ and $10\mu\text{m}$ wavelengths and we obtain $\beta_{\text{ext}}(4\mu\text{m}) \approx 4.9$

$\beta_{\text{ext}}(10\mu\text{m})$, in good agreement with the exact Mie results in Figs. 9A and 9B. At the $0.55\mu\text{m}$ wavelength the Chýlek approximation cannot legitimately be used since particles with radii greater than the maximum value ($r_m \approx 0.5\mu\text{m}$) contribute a significant part of the extinction for some haze distributions. Nevertheless, if we apply the approximation we obtain $\beta_{\text{ext}}(0.55\mu\text{m}) \approx 6.5 \beta_{\text{ext}}(4\mu\text{m})$, a result that is within a factor 2 of the exact Mie results in Figs. 9A and 9B.

Since the reader may be interested in the radiative effects of fog and haze particles at other wavelengths in the infrared atmospheric windows, we have included in the appendix calculations of extinction vs LWC (for the 289 distributions measured at Grafenwöhr) at $\lambda = 3\mu\text{m}$, $5\mu\text{m}$, $8\mu\text{m}$, $11\mu\text{m}$, and $12\mu\text{m}$ (Fig. A6). These data are replotted in the form of extinction at $\lambda = 0.55\mu\text{m}$ vs extinction at $\lambda = 1.2\mu\text{m}$, $3\mu\text{m}$, $4\mu\text{m}$, $5\mu\text{m}$, $8\mu\text{m}$, $10\mu\text{m}$, $11\mu\text{m}$, and $12\mu\text{m}$ in Fig. A7. Approximate empirical relationships between extinction at a particular infrared wavelength and extinction at $\lambda = 0.55\mu\text{m}$ can be derived from Fig. A7. For example, fitting the points in Fig. A7-c and A7-f with straight lines yields:

$$\beta_{\text{ext}}(4\mu\text{m}) \approx 0.12 \beta_{\text{ext}}(0.55\mu\text{m})^{1.75},$$

and

$$\beta_{\text{ext}}(10\mu\text{m}) \approx 0.036 \beta_{\text{ext}}(0.55\mu\text{m})^{1.72}.$$

CONCLUSIONS

Measurements of haze and fog droplet distributions carried out during 8 days in wintertime in West Germany show little vertical structure for haze, but a general trend toward increasing concentration of $4\mu\text{m}$ to $16\mu\text{m}$ radius droplets with altitude for fog conditions with visibilities < 1 km. Particulate extinction calculated from measured distributions at $0.55\mu\text{m}$, $4\mu\text{m}$, and $10\mu\text{m}$ wavelengths shows increases by a factor of 2 to 1000 over the first 150 m altitude for fog. Extinction is found to be approximately proportional to $1/\lambda$ over this spectral region for haze conditions, but nearly independent of wavelength for fog. Thus, for fog, from the point of view of extinction of radiation, there is no advantage of an infrared imaging system operating in the $8\text{--}12\mu\text{m}$ atmospheric window over one operating in the $3\text{--}5\mu\text{m}$ window. There exists a size distribution independent linear relationship between particle extinction coefficient and liquid water content at $10\mu\text{m}$. However, the approximate relationships found between extinction (from visible through

4 μ m wavelengths) and liquid water content are attributed to a common form of the fog and haze size distributions measured.

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APPENDIX FIGURES

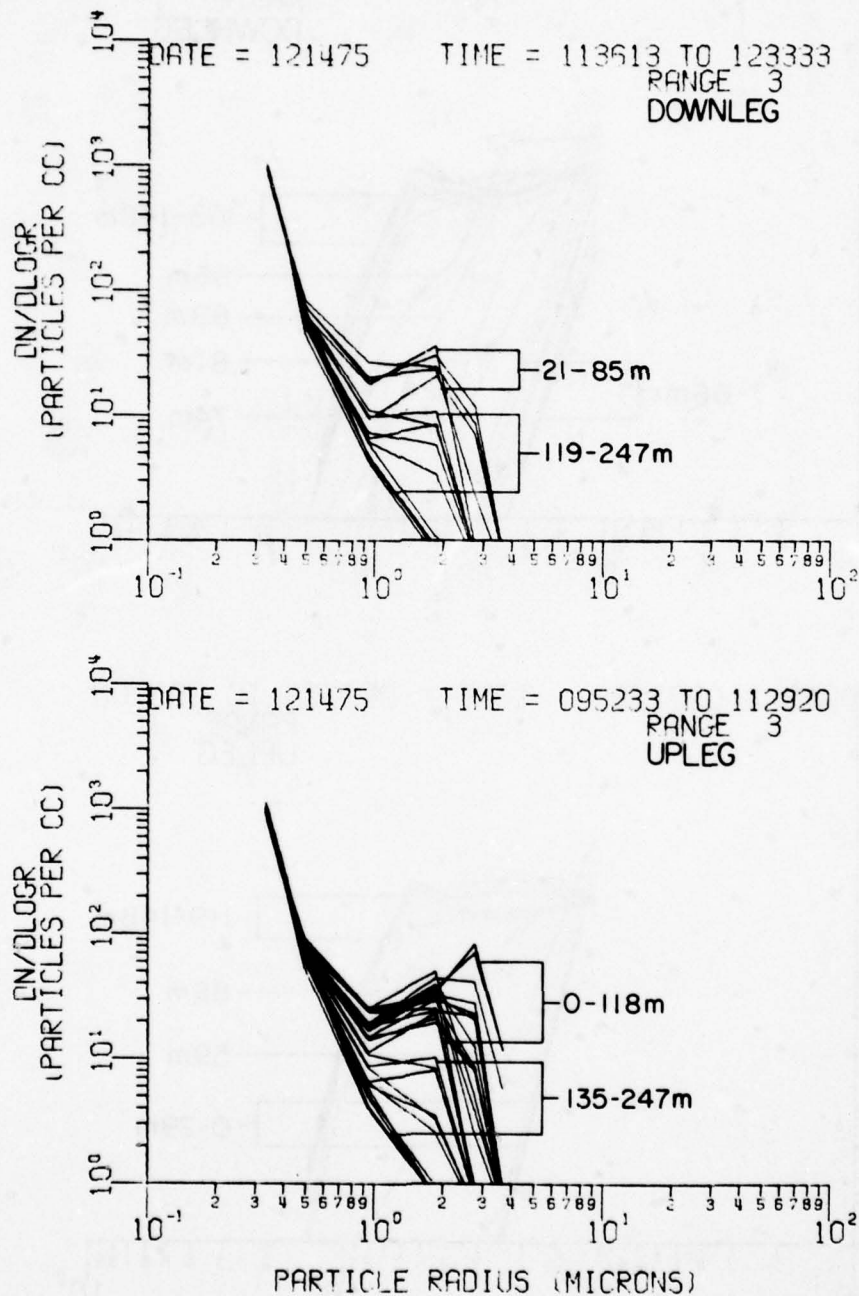


Figure A1-a Particle number size distribution measurements made with a tethered balloon-borne aerosol counter for 20 upleg and downleg profiles during haze and fog conditions. The date, local time (hours, minutes, seconds), and time interval for the measurements are given, as well as the size range setting (see Fig. 4) of the aerosol counter.

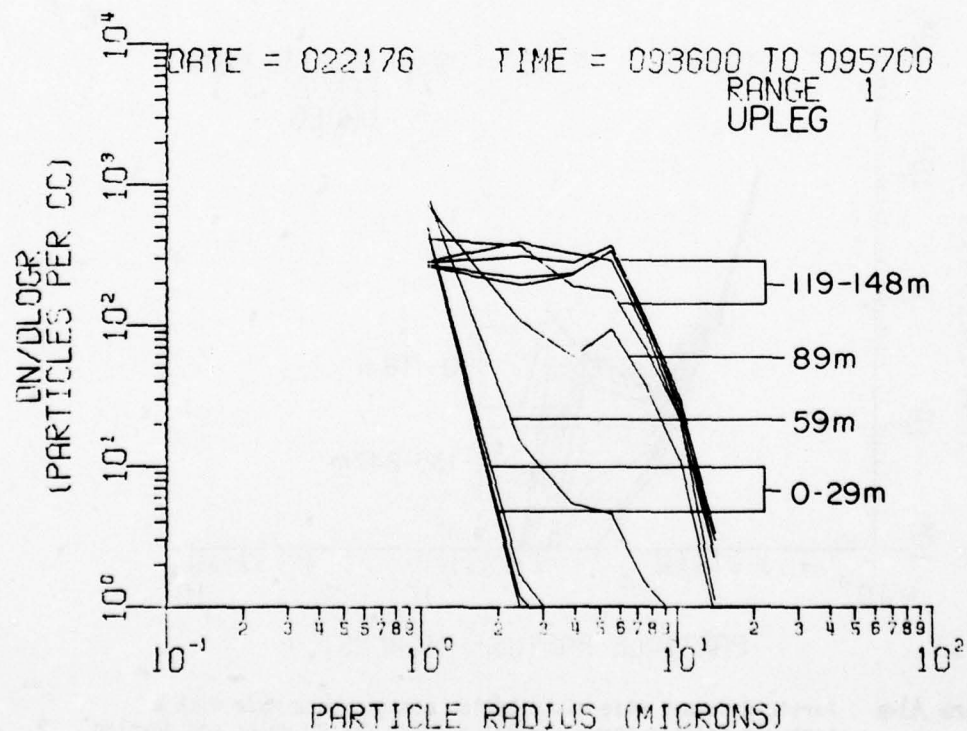
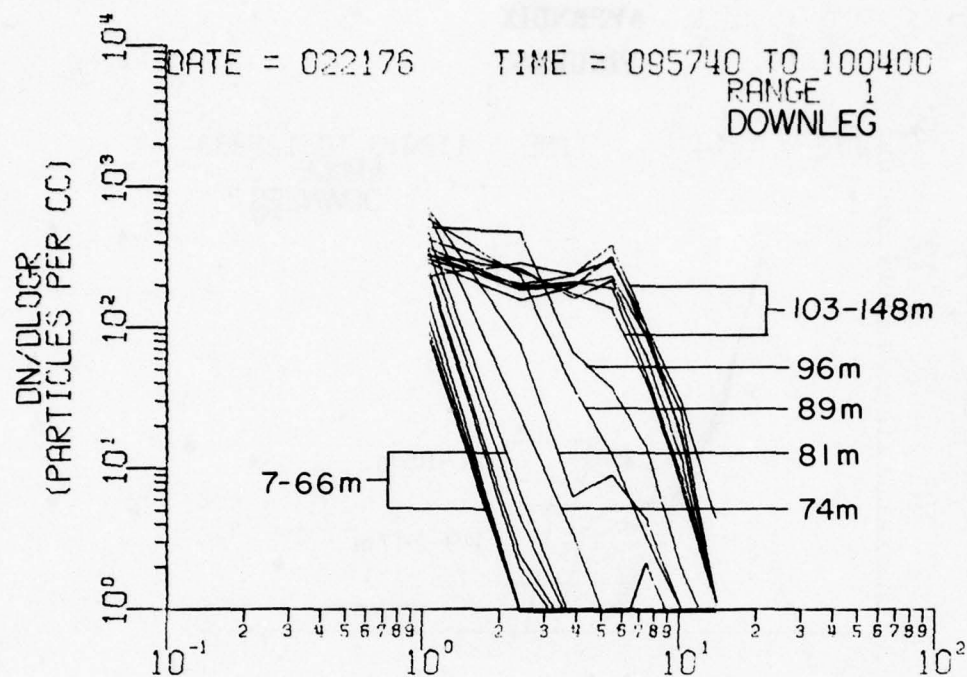


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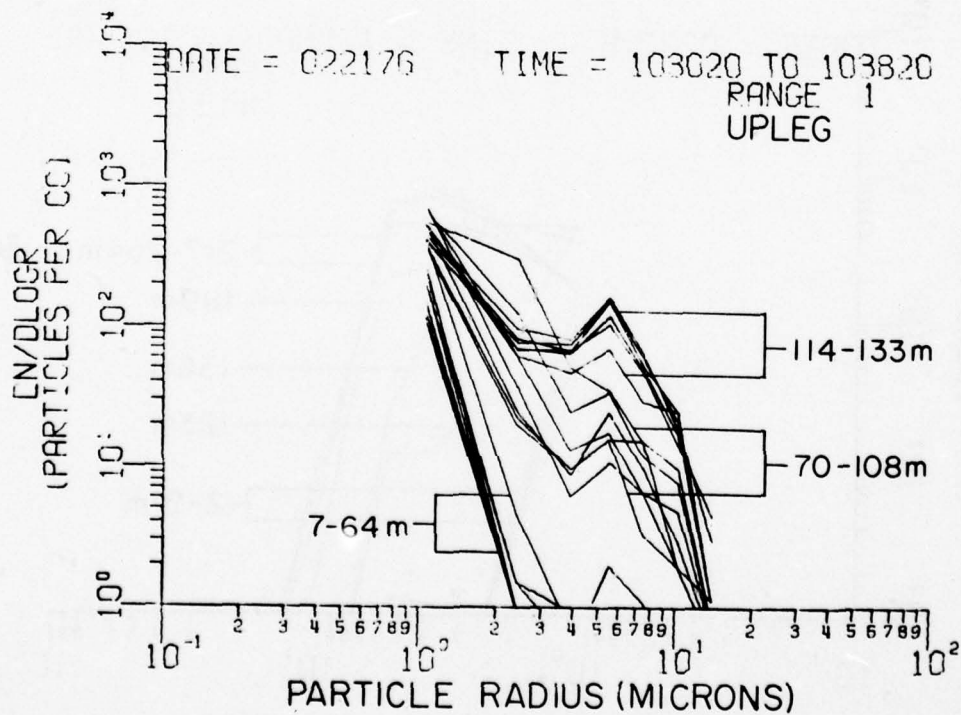
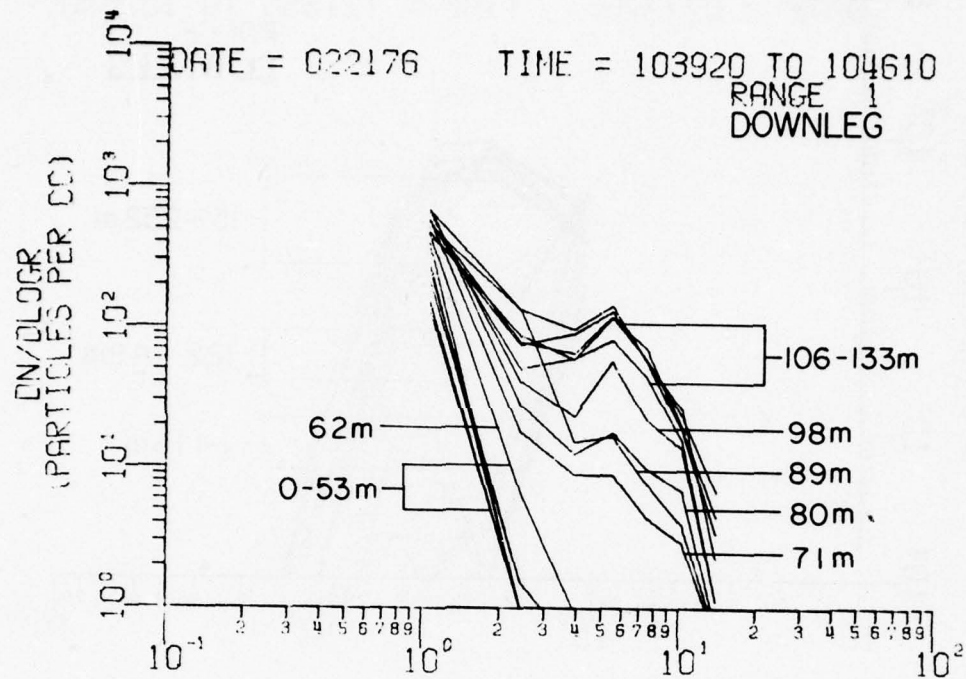


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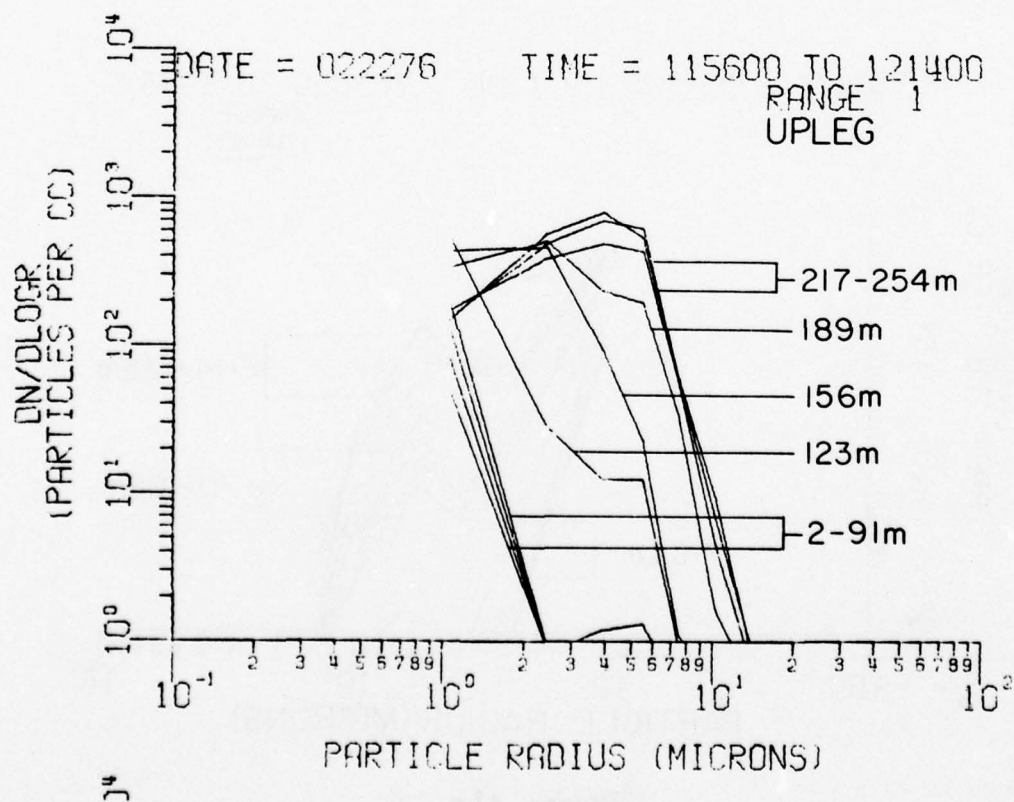
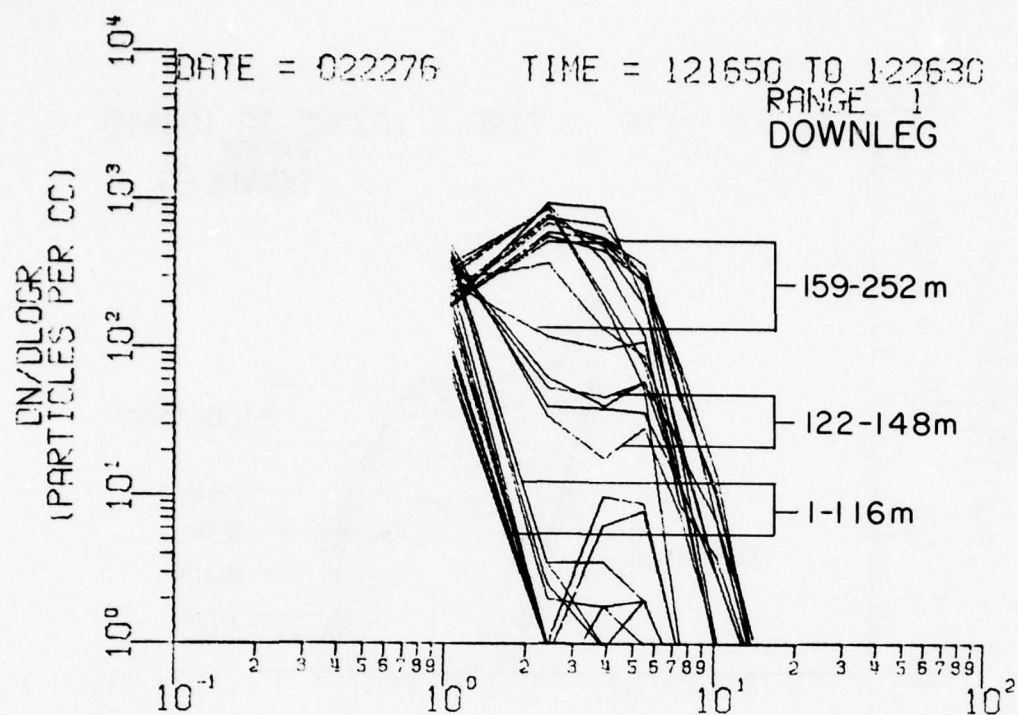


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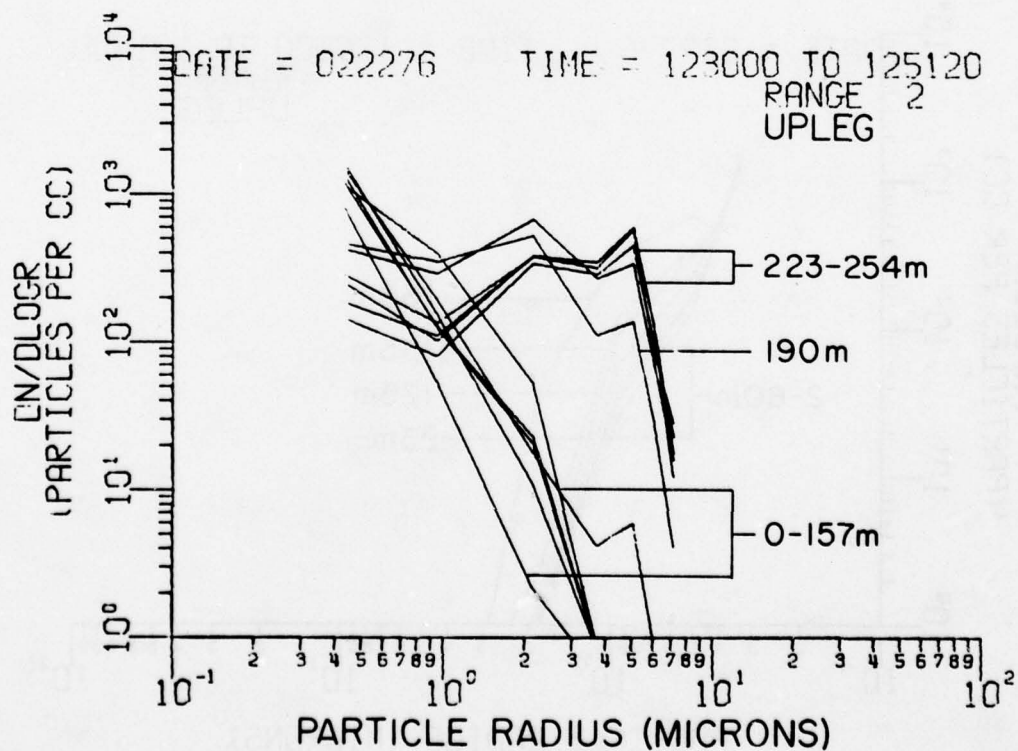
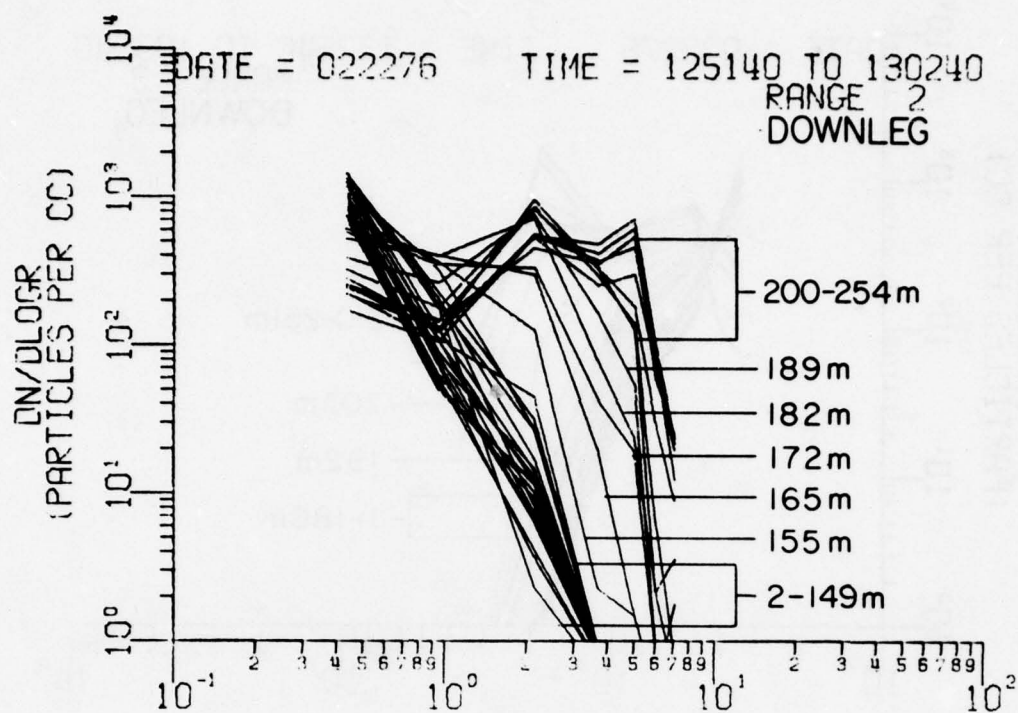


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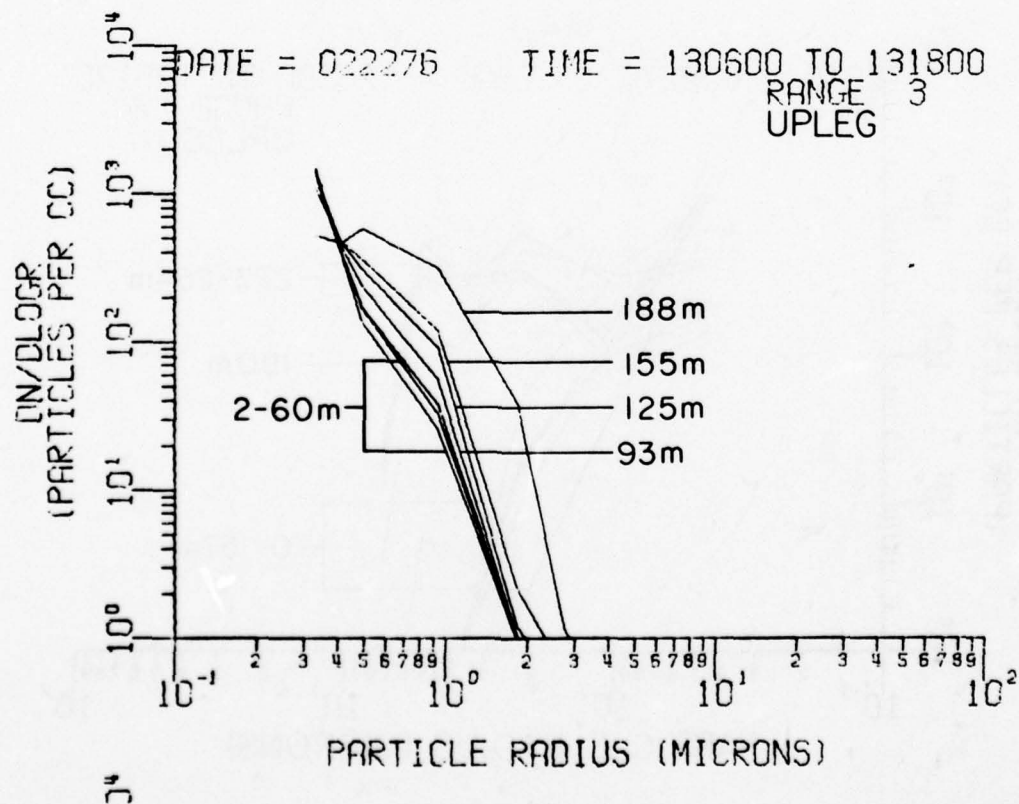
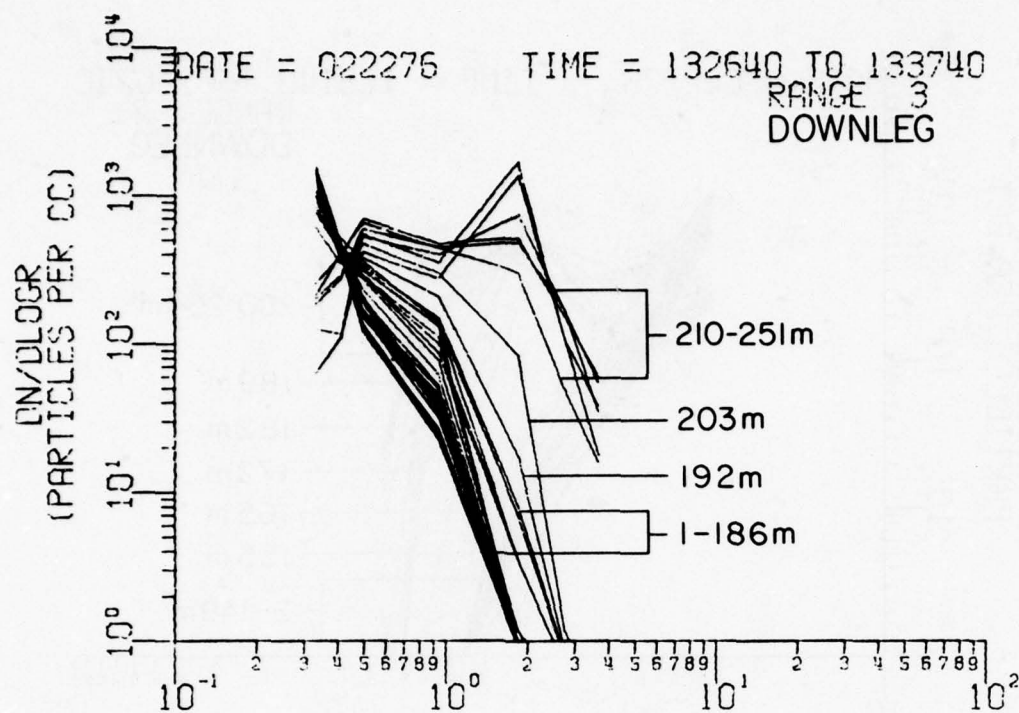


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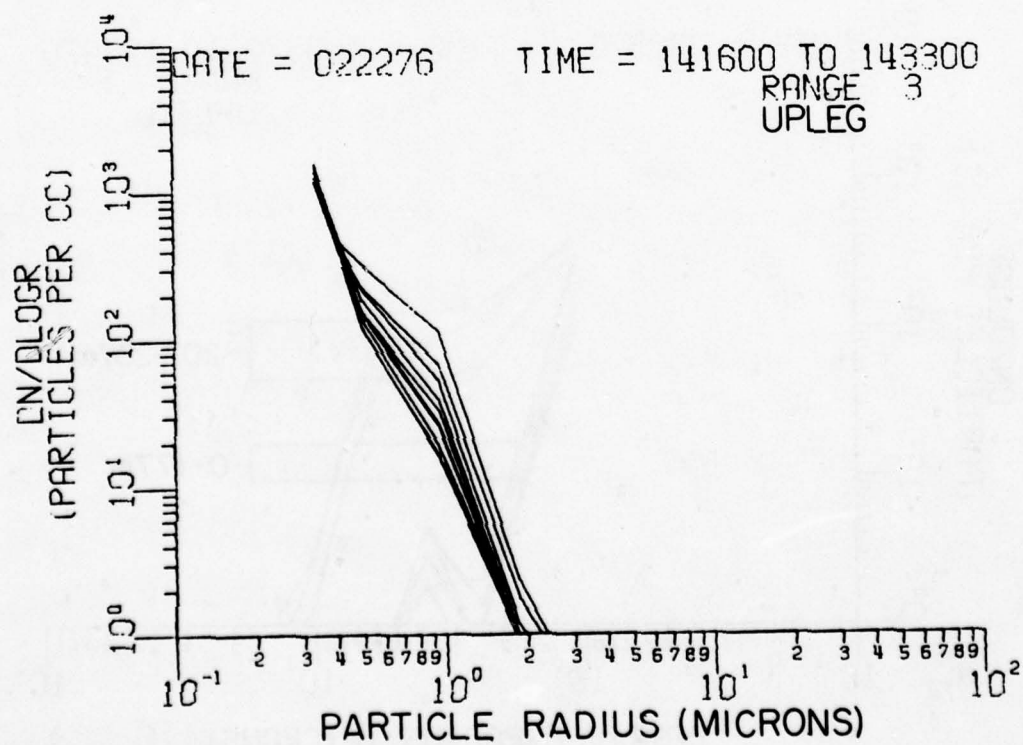
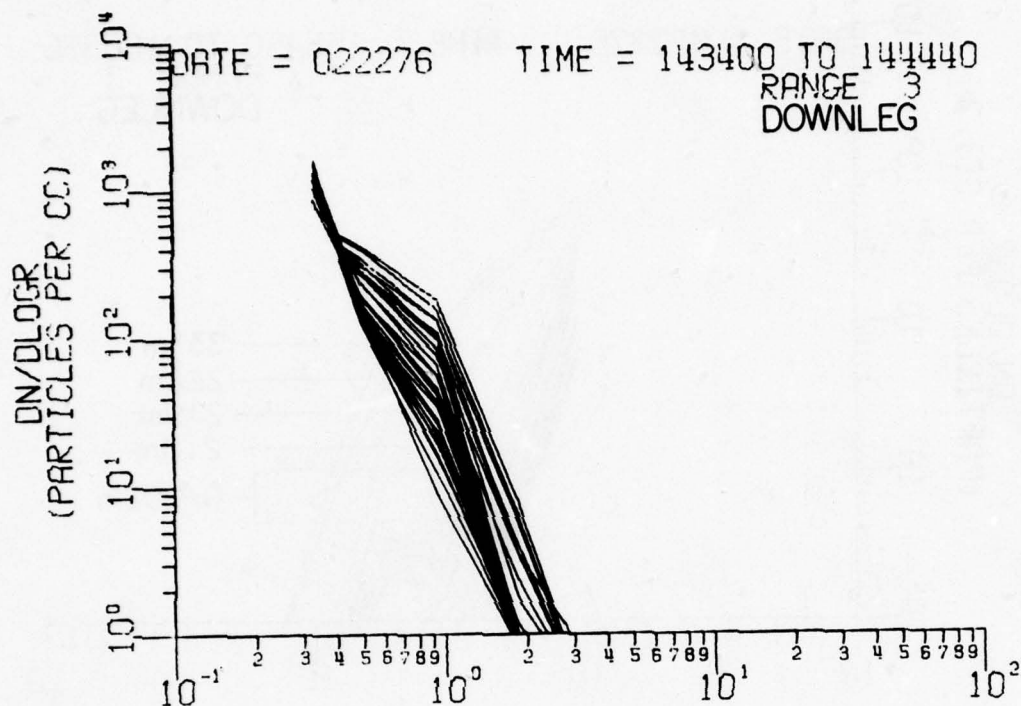


Figure A1-g

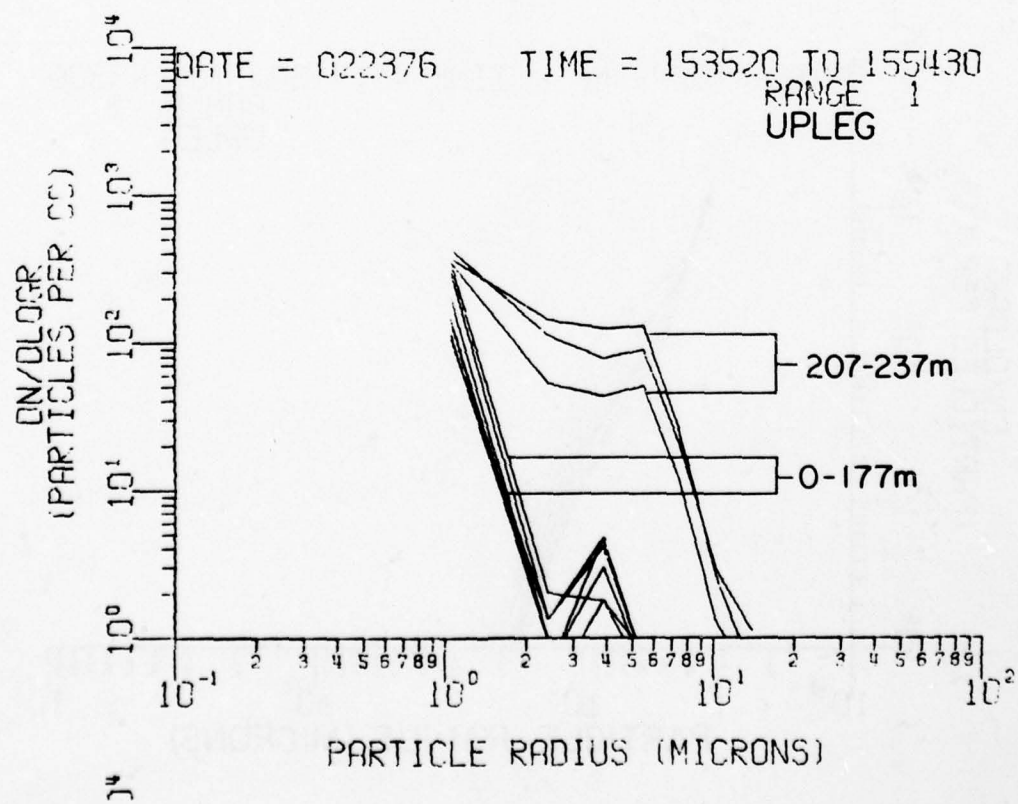
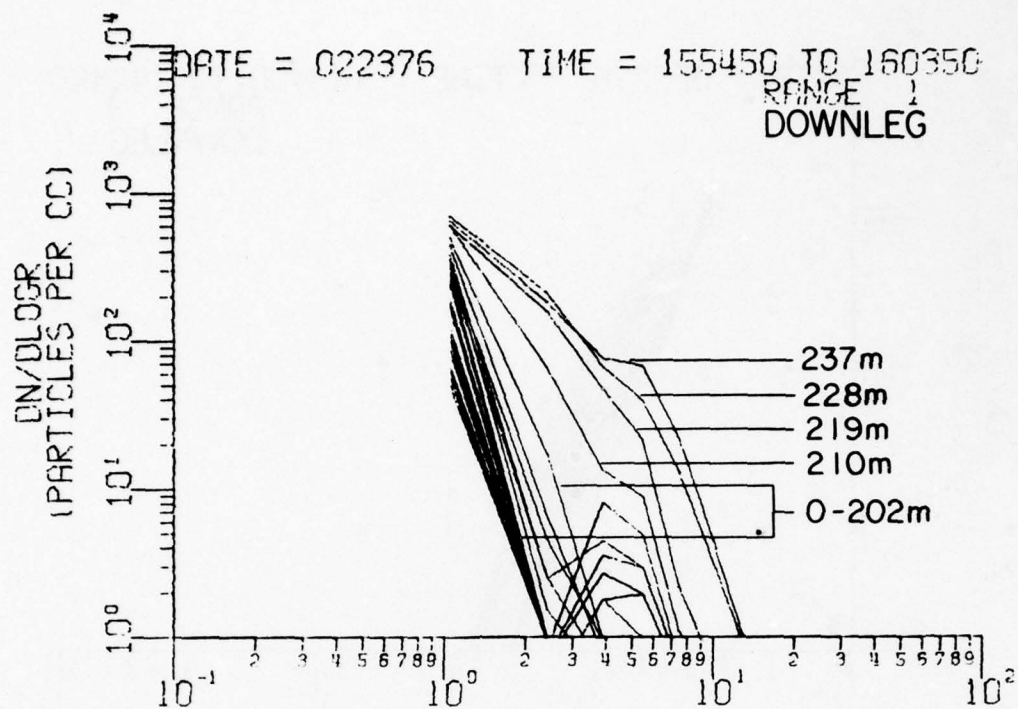


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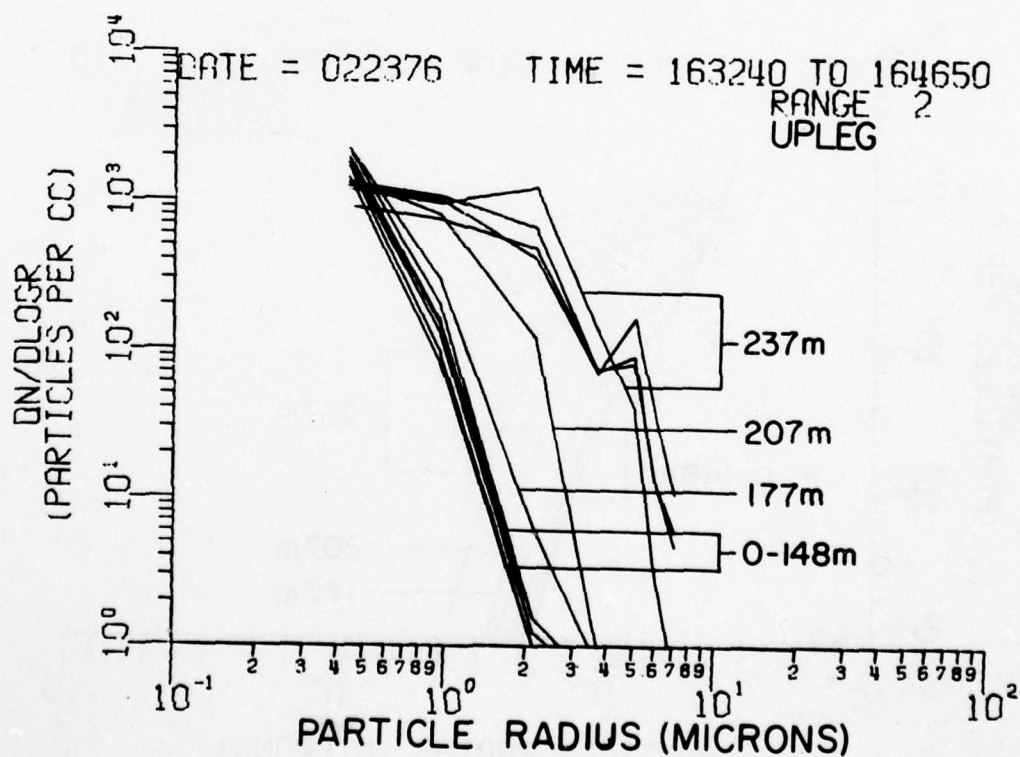
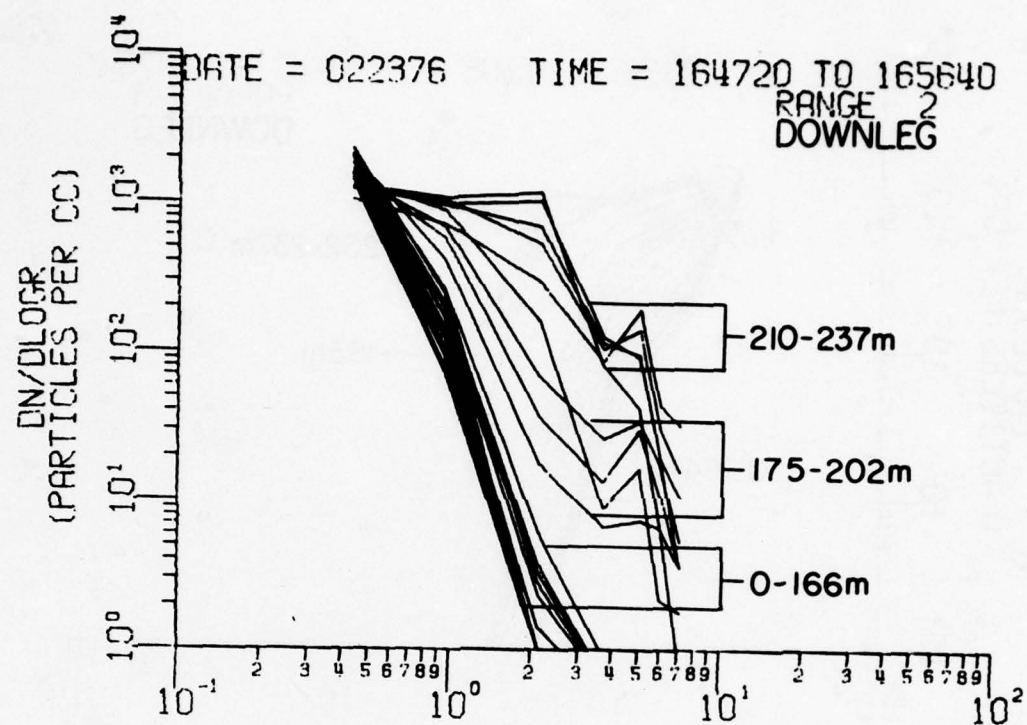


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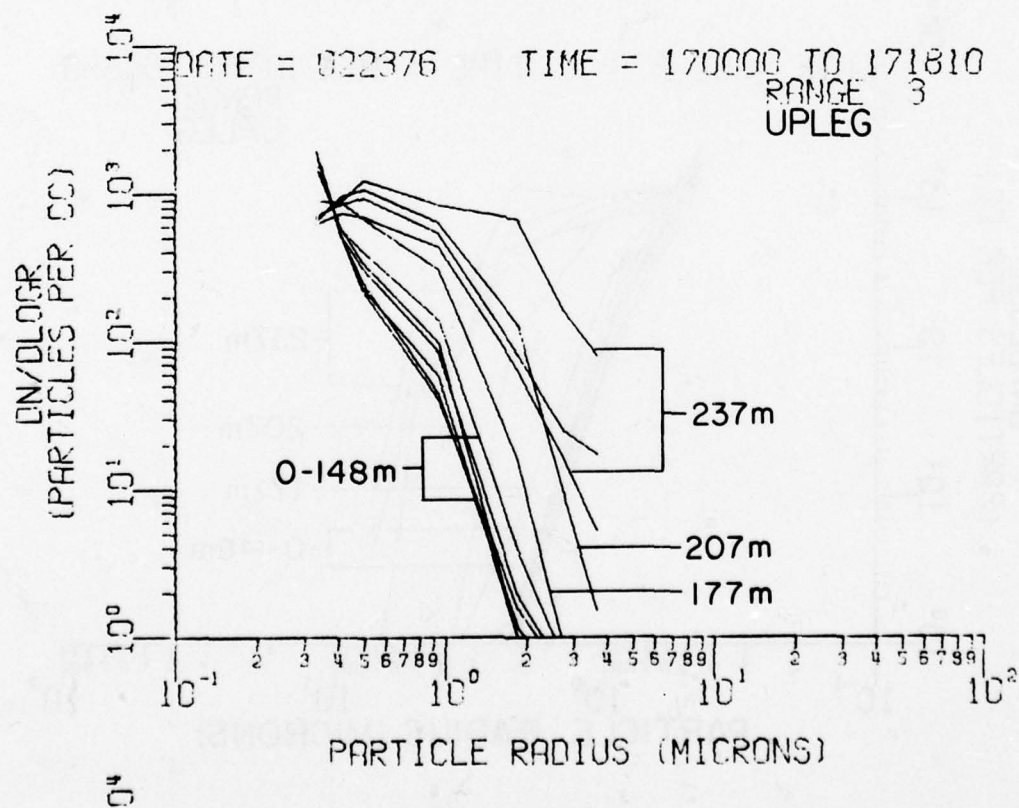
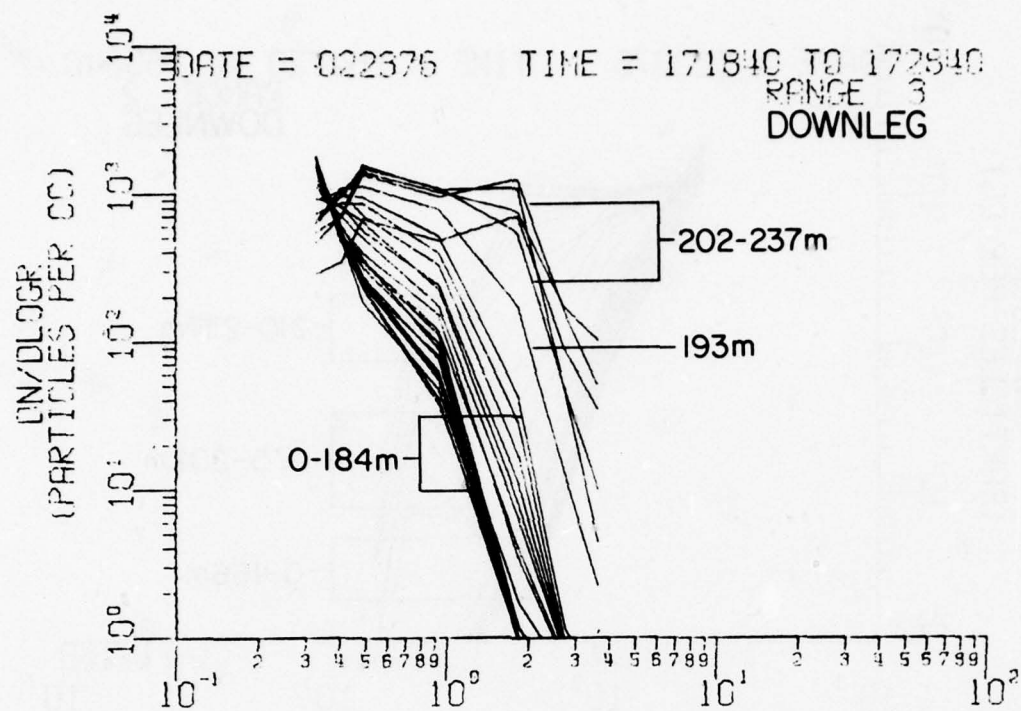


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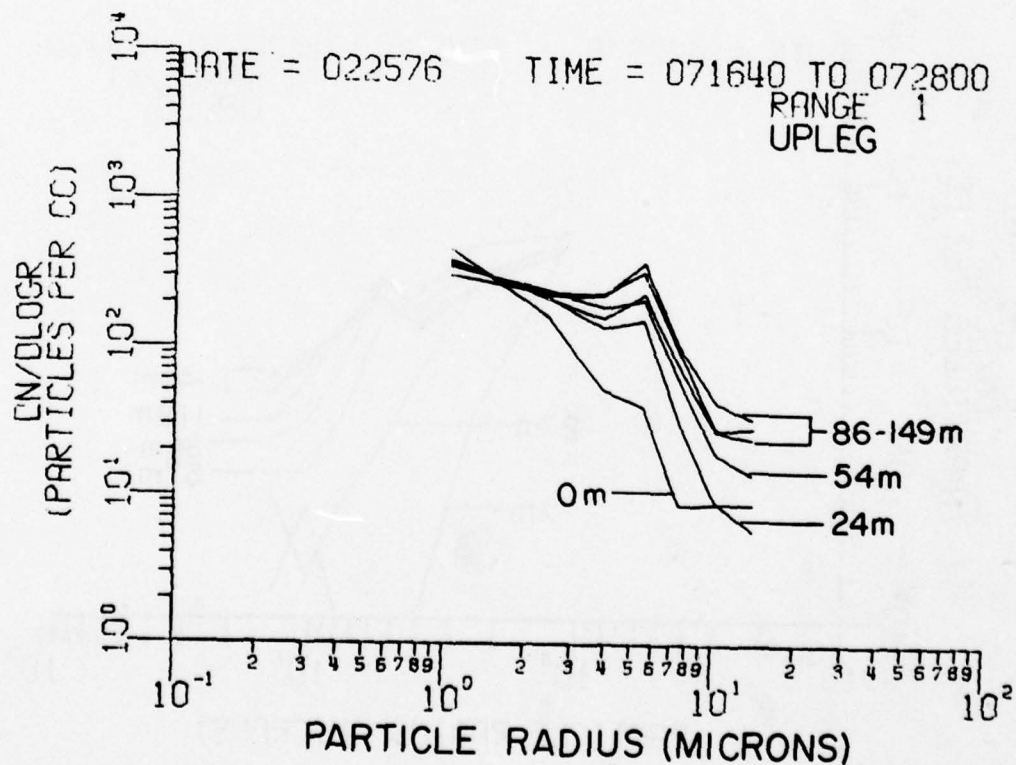
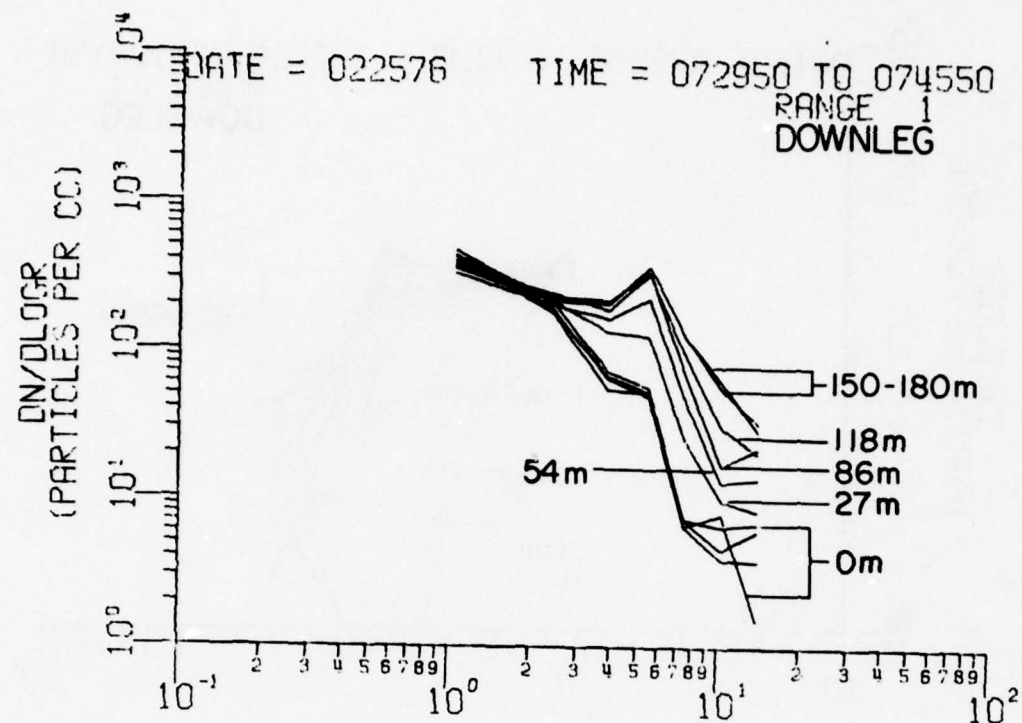


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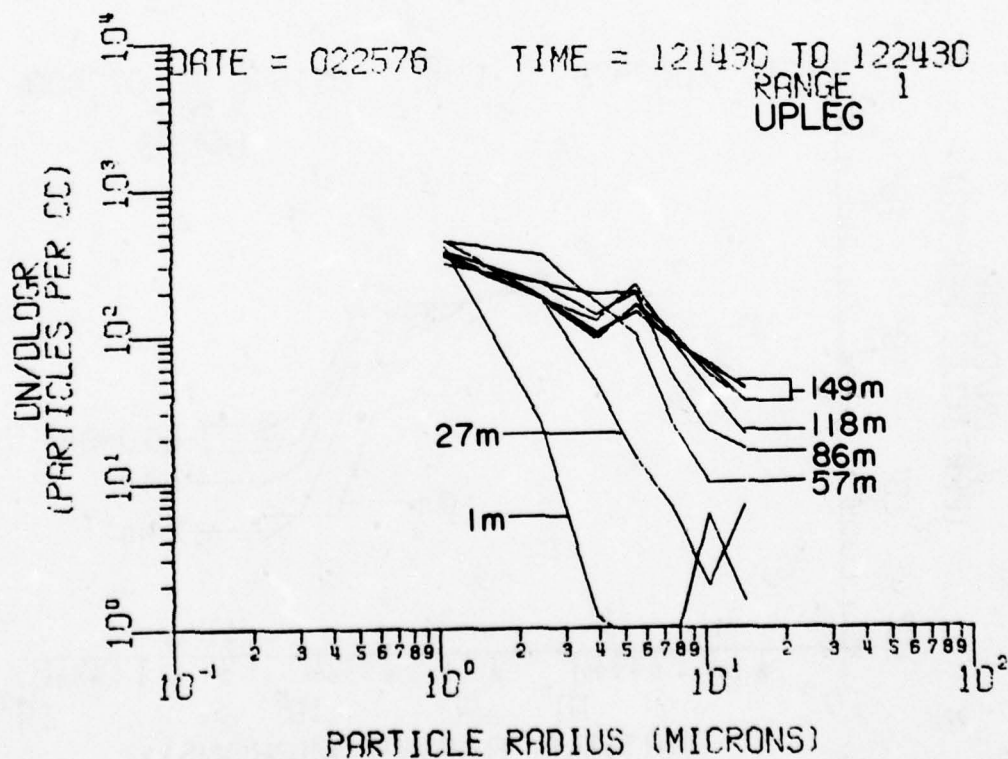
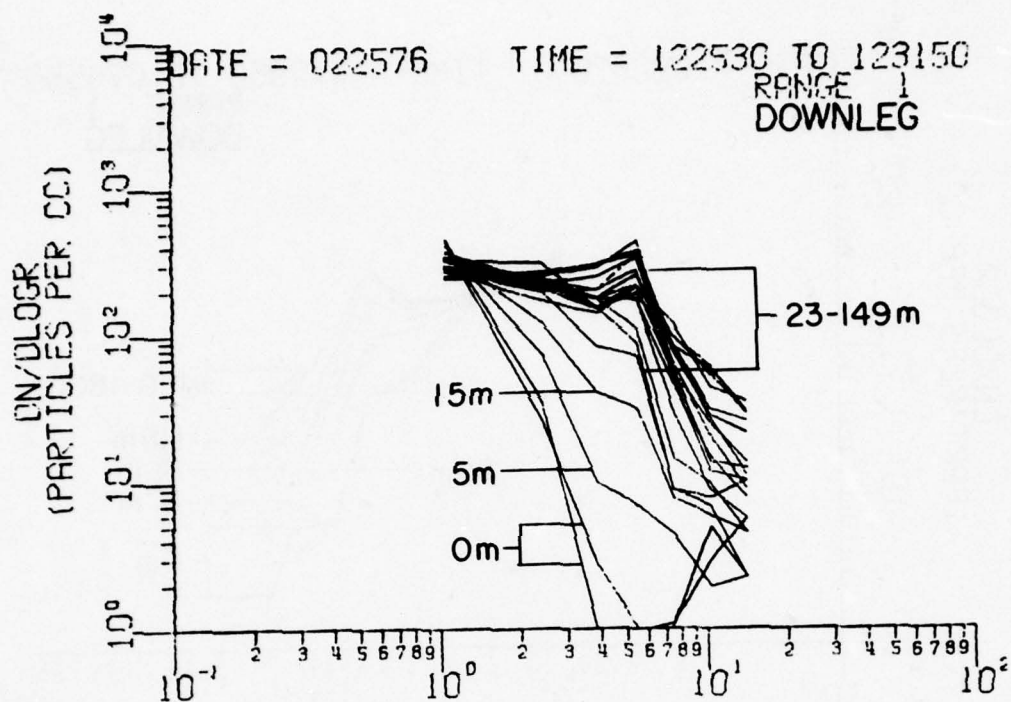


Figure A1-1

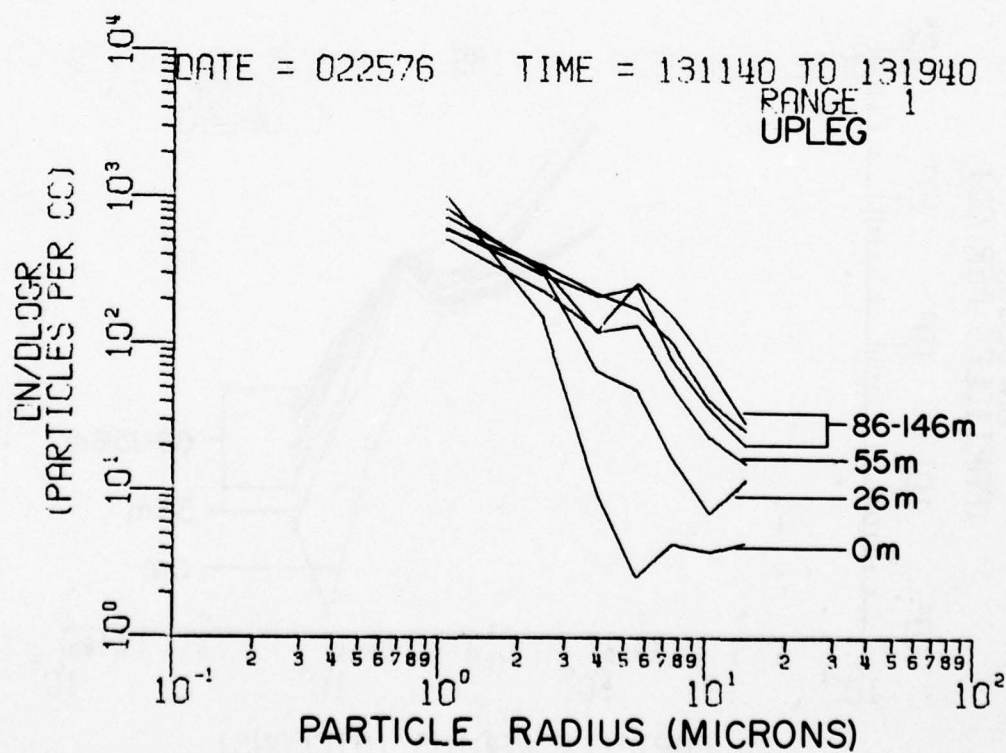
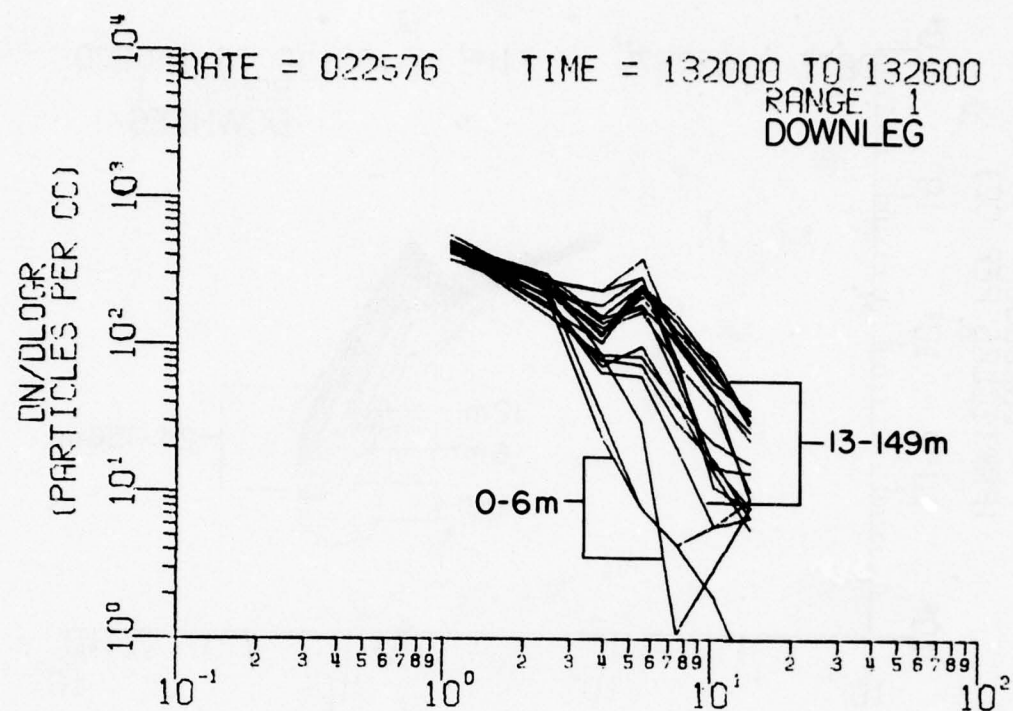


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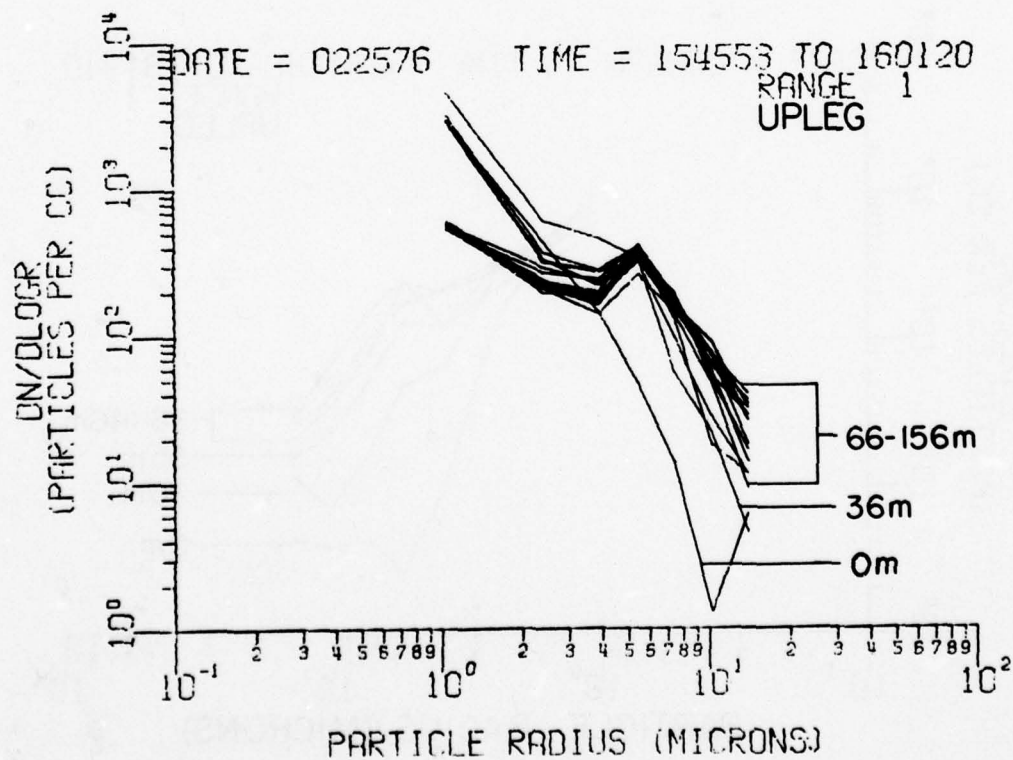
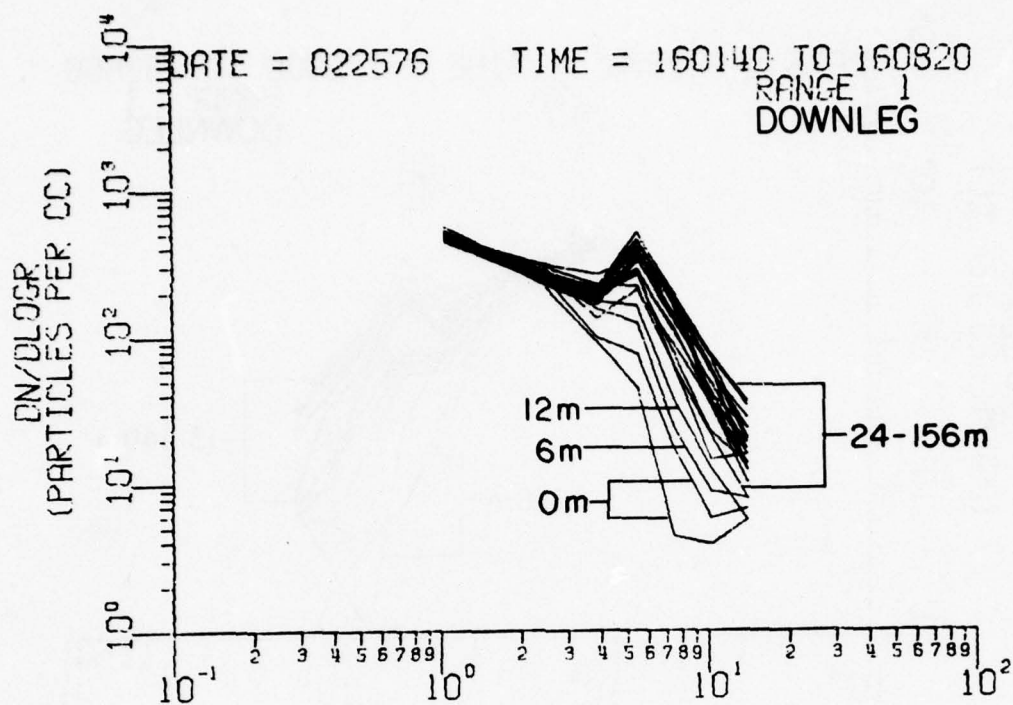


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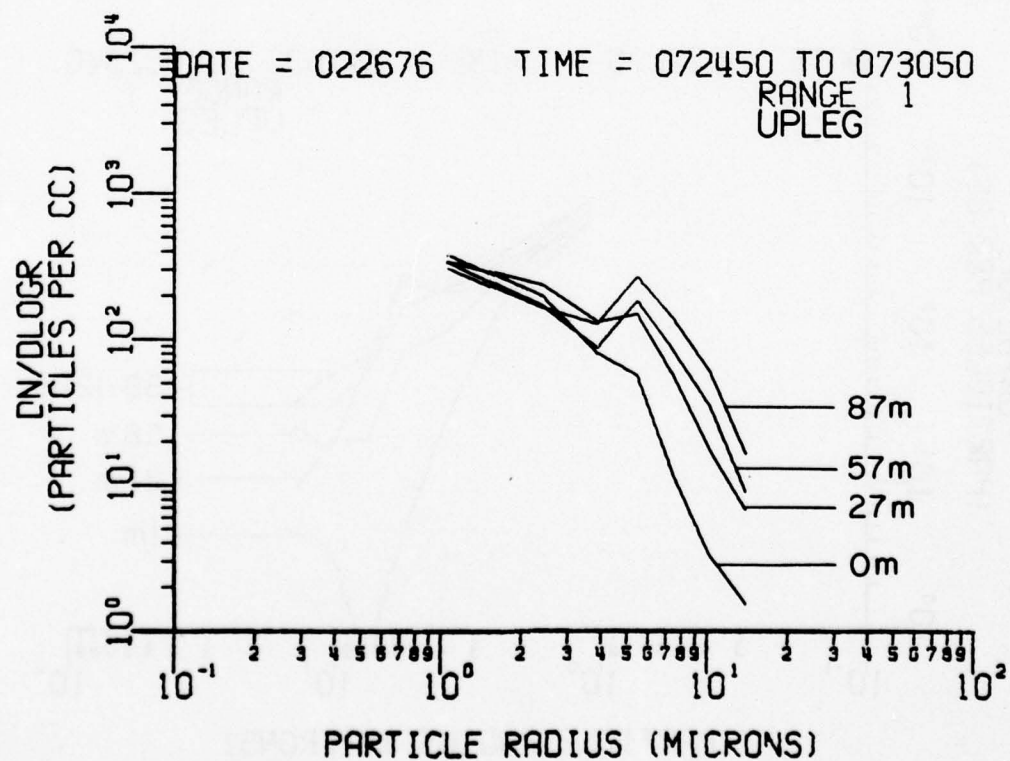
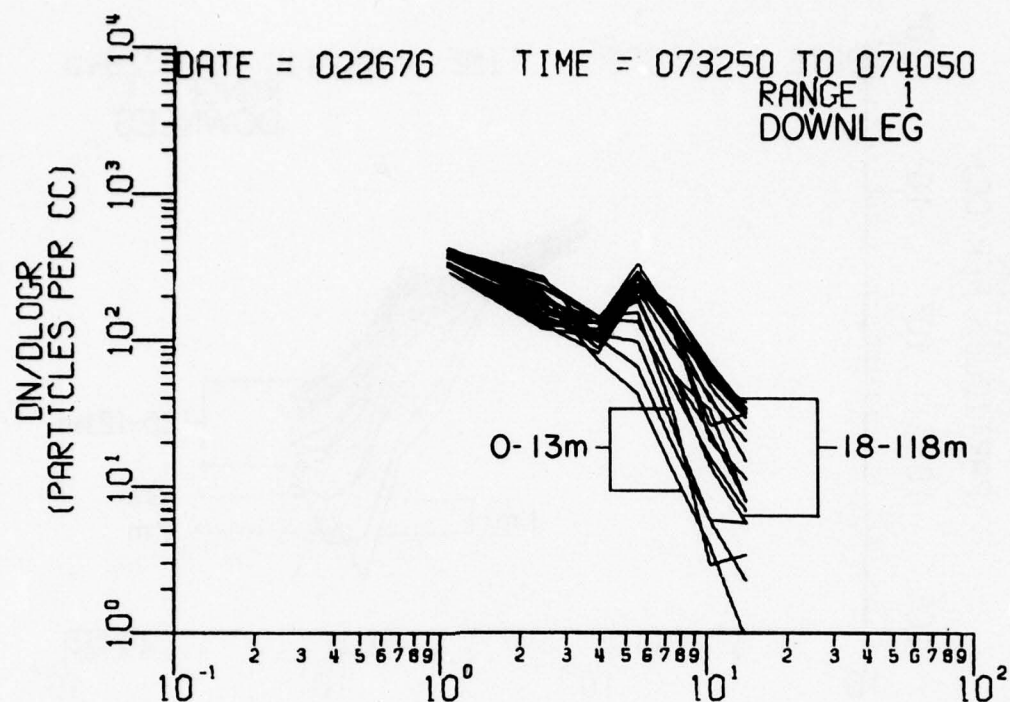


Figure A1-o

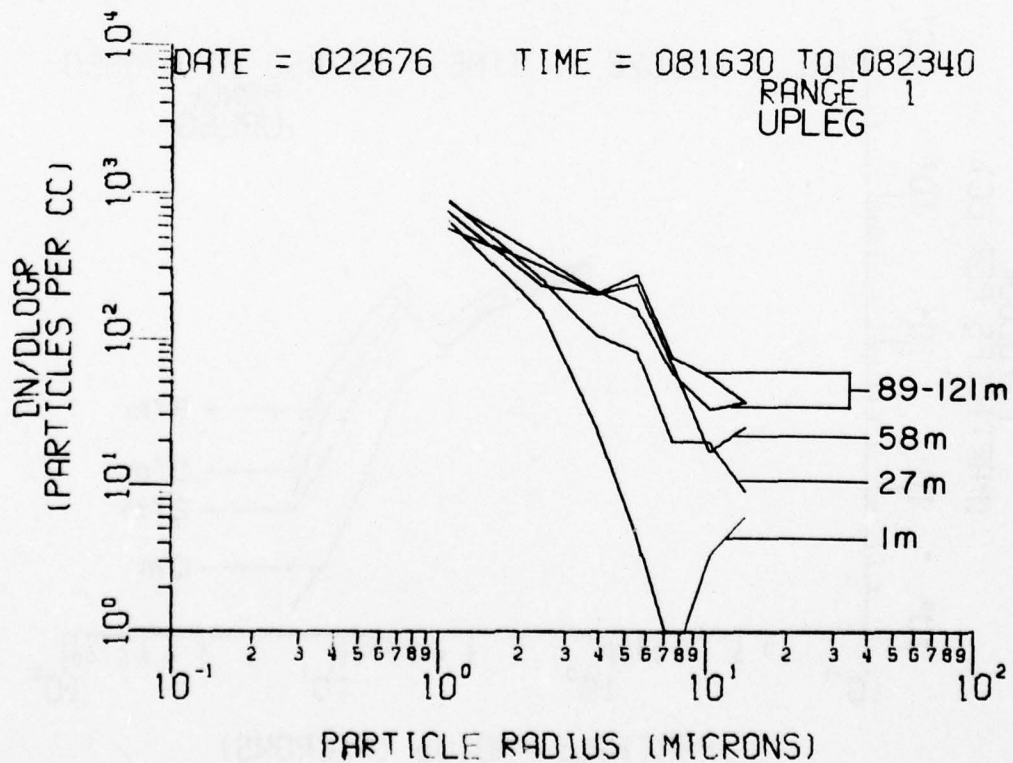
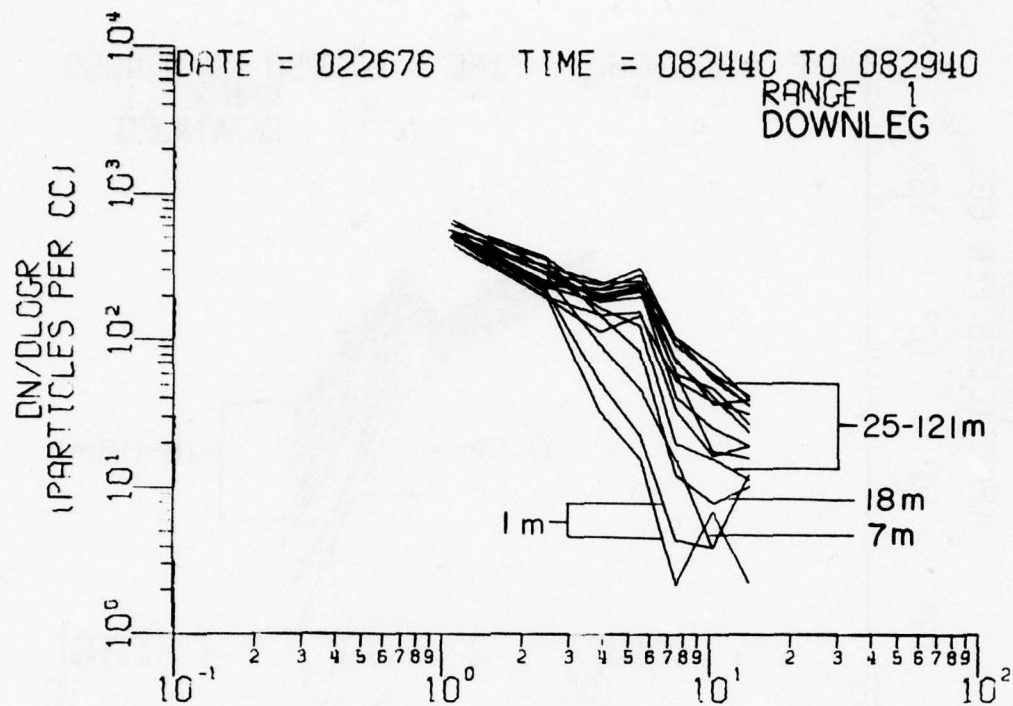


Figure A1-p

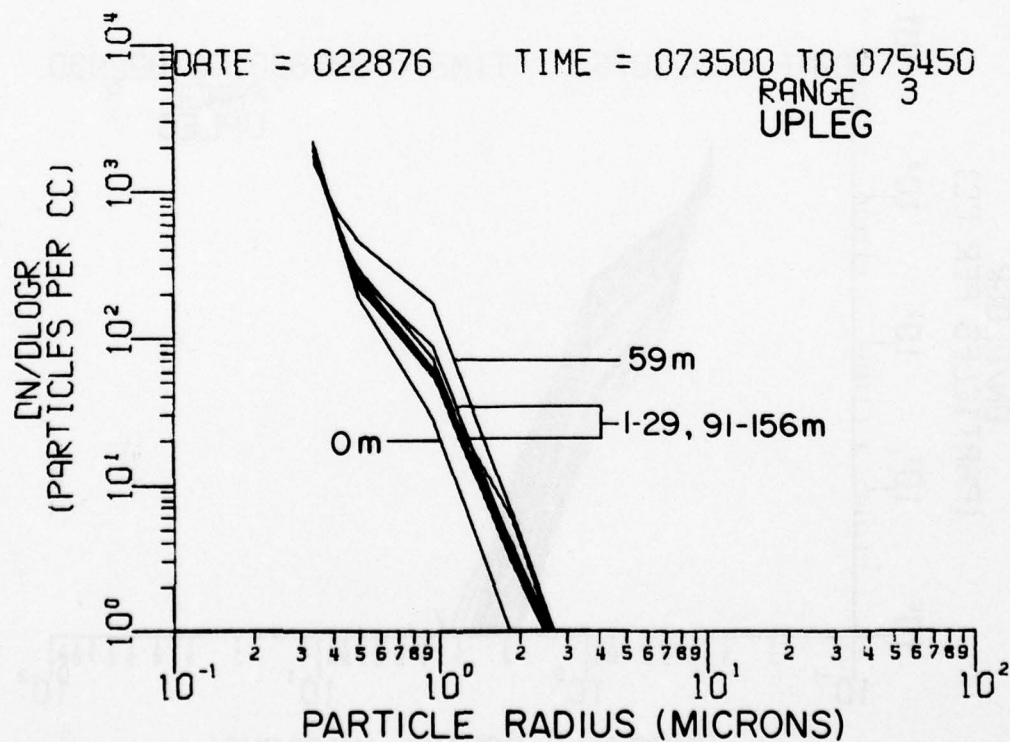
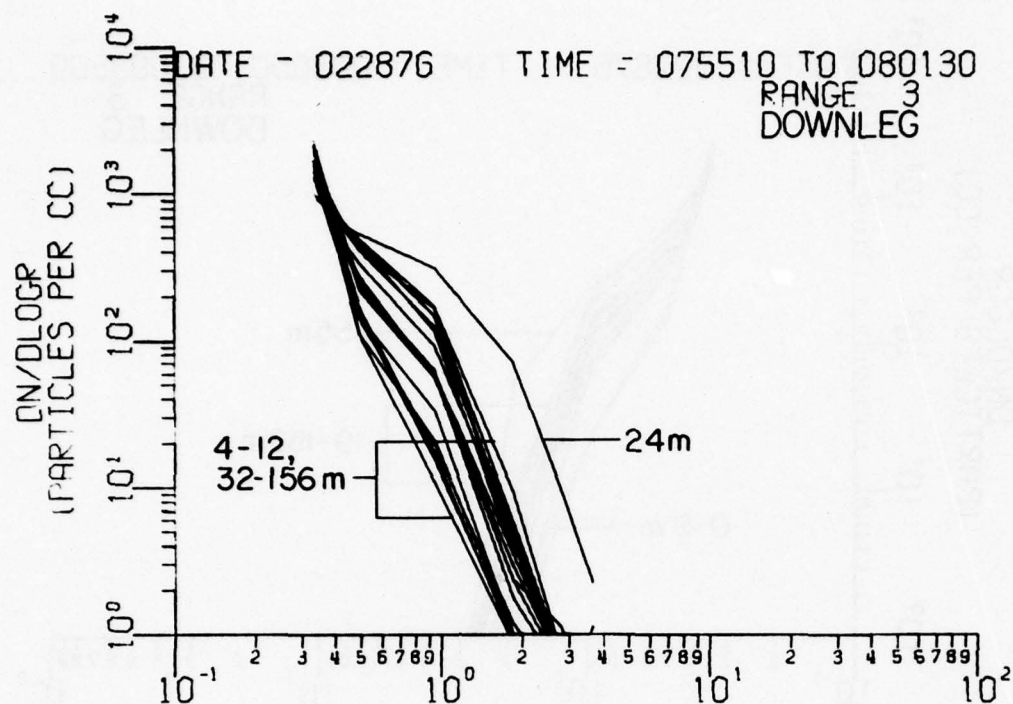


Figure A1-q

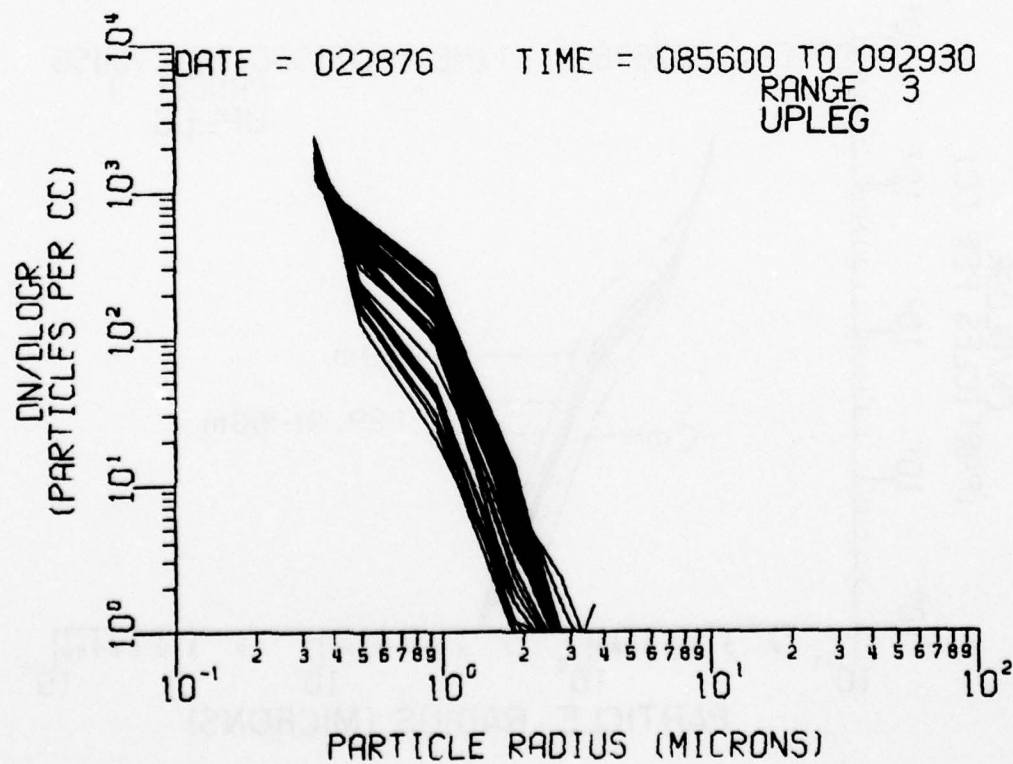
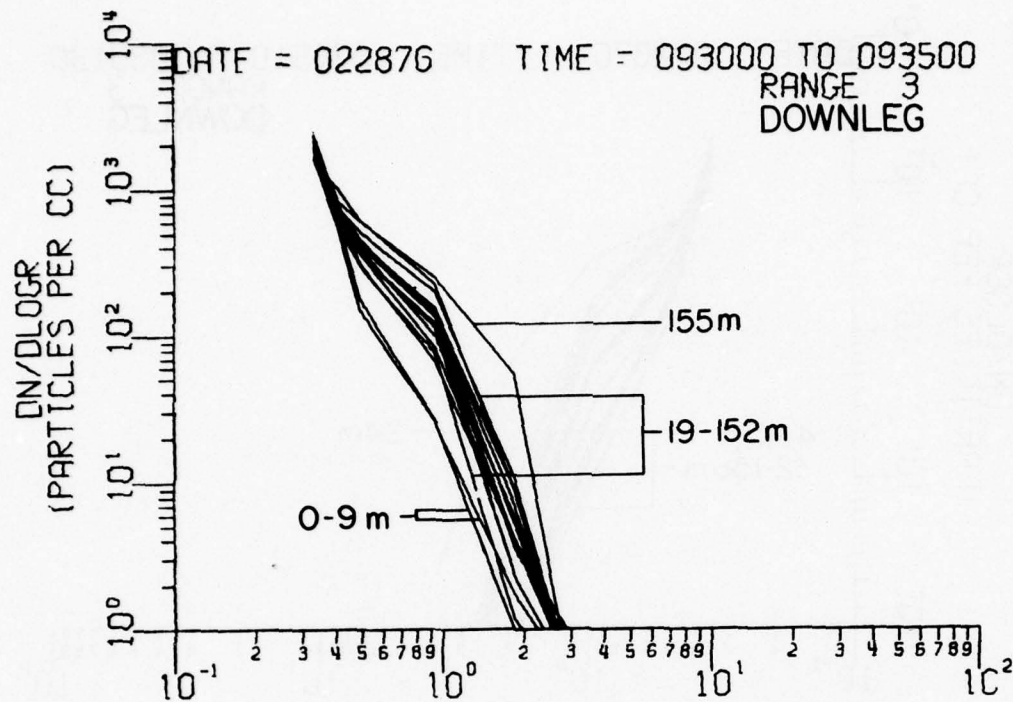


Figure A1-r

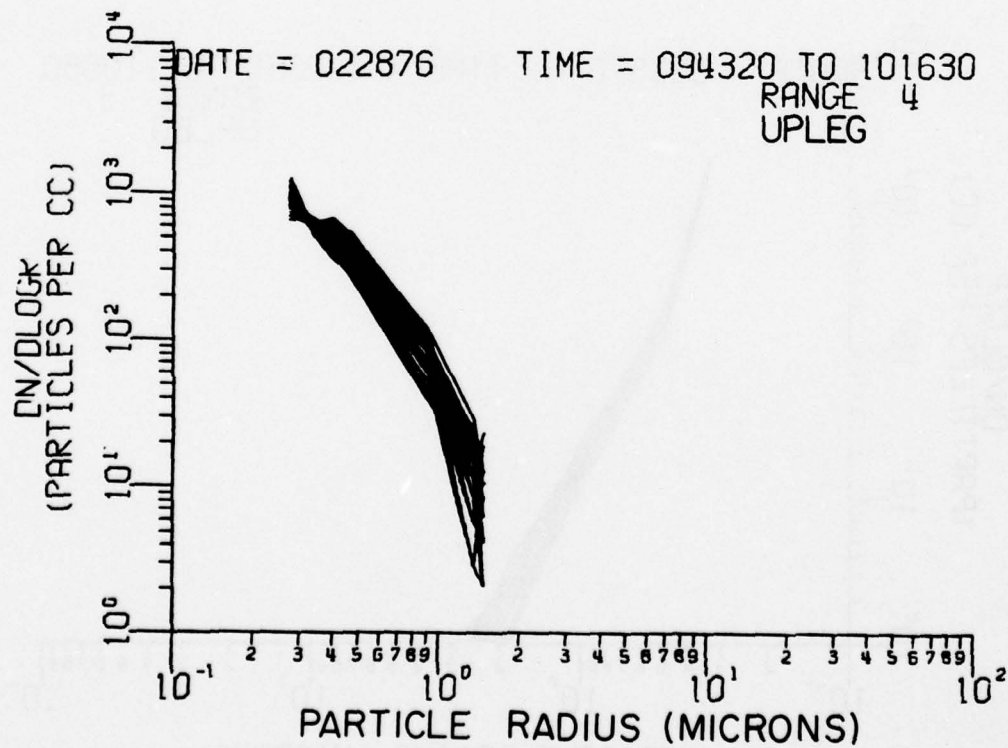
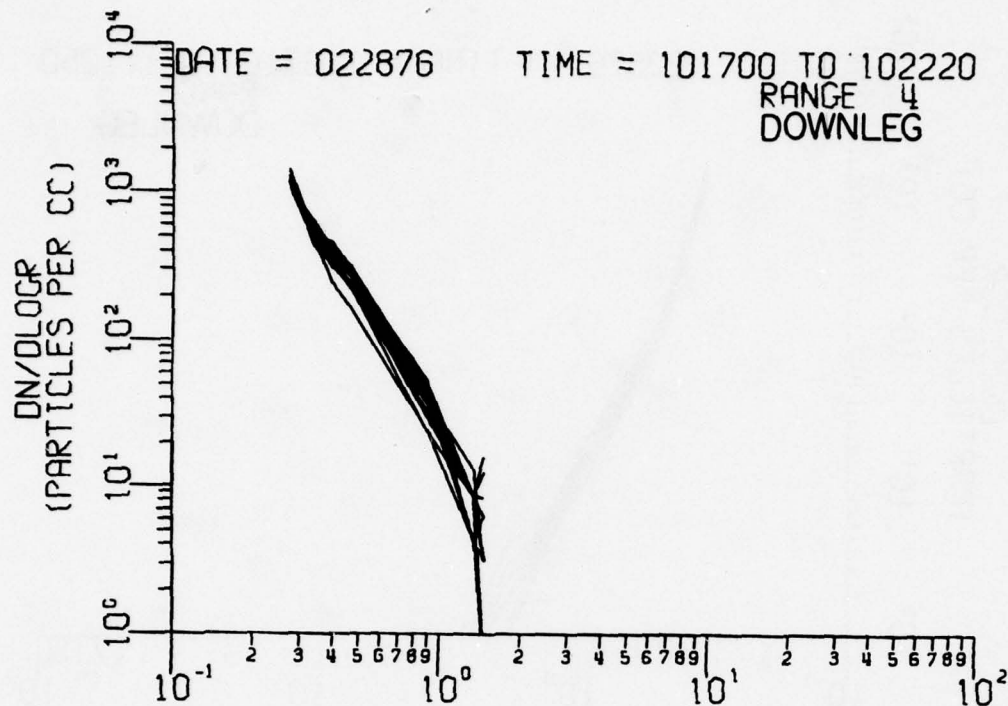


Figure A1-a

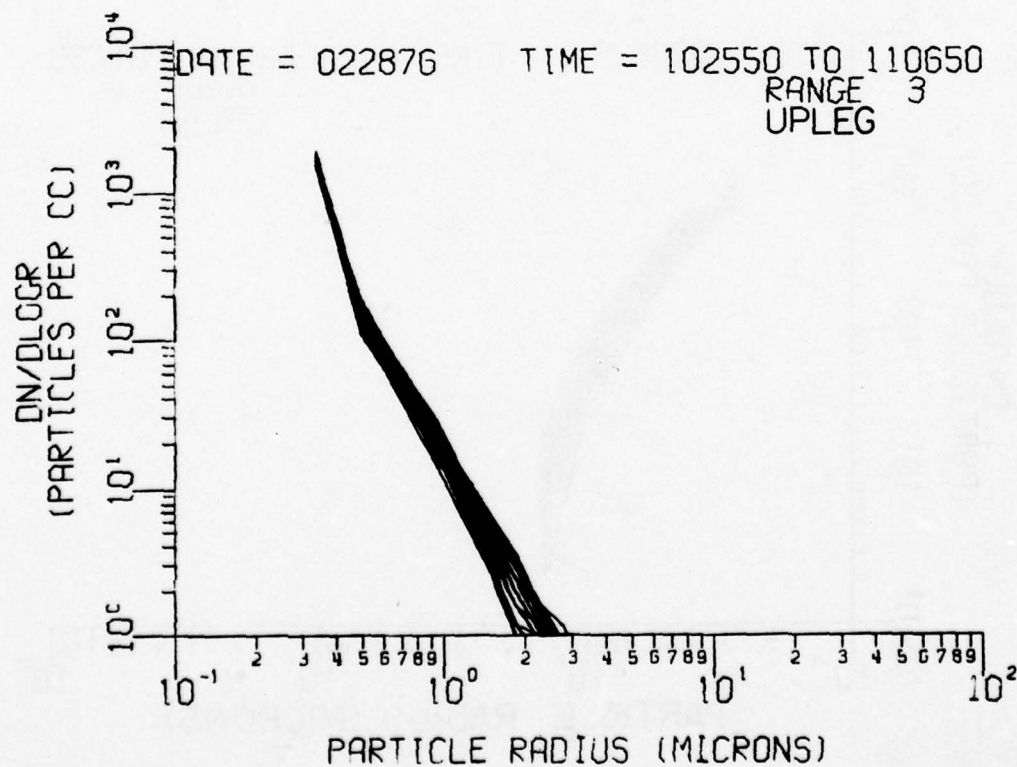
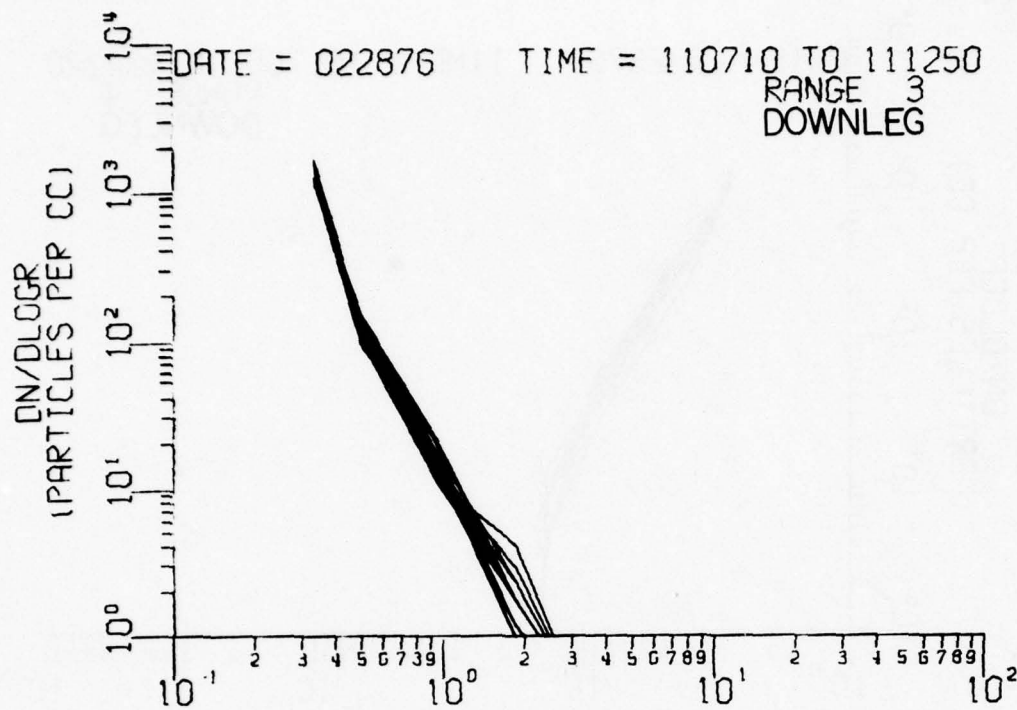


Figure Al-t

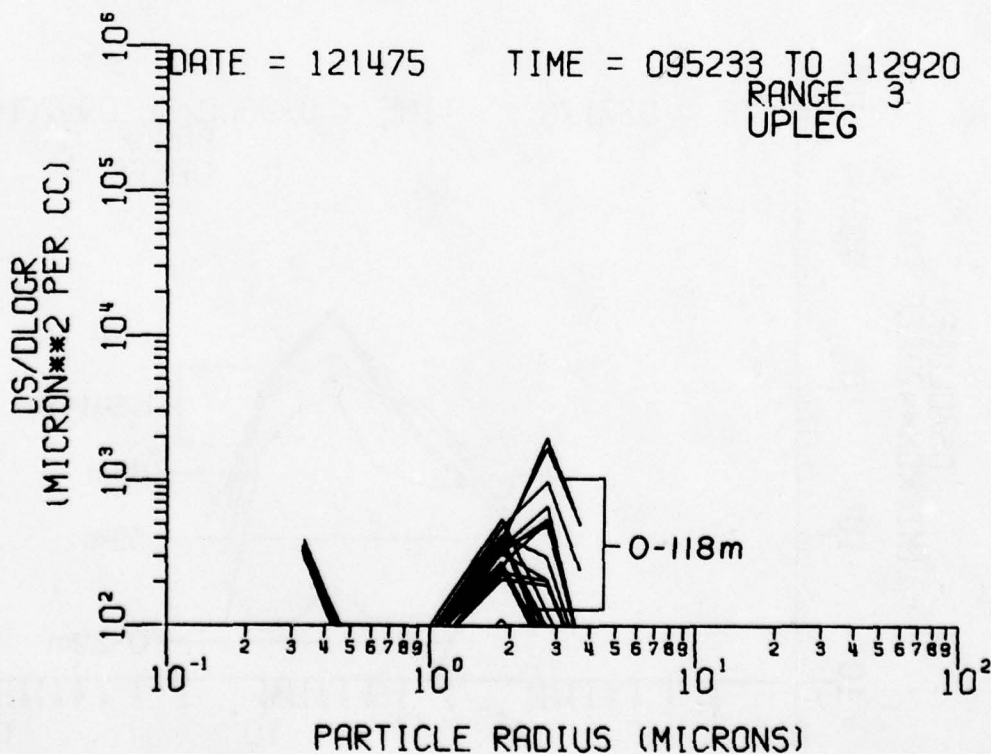
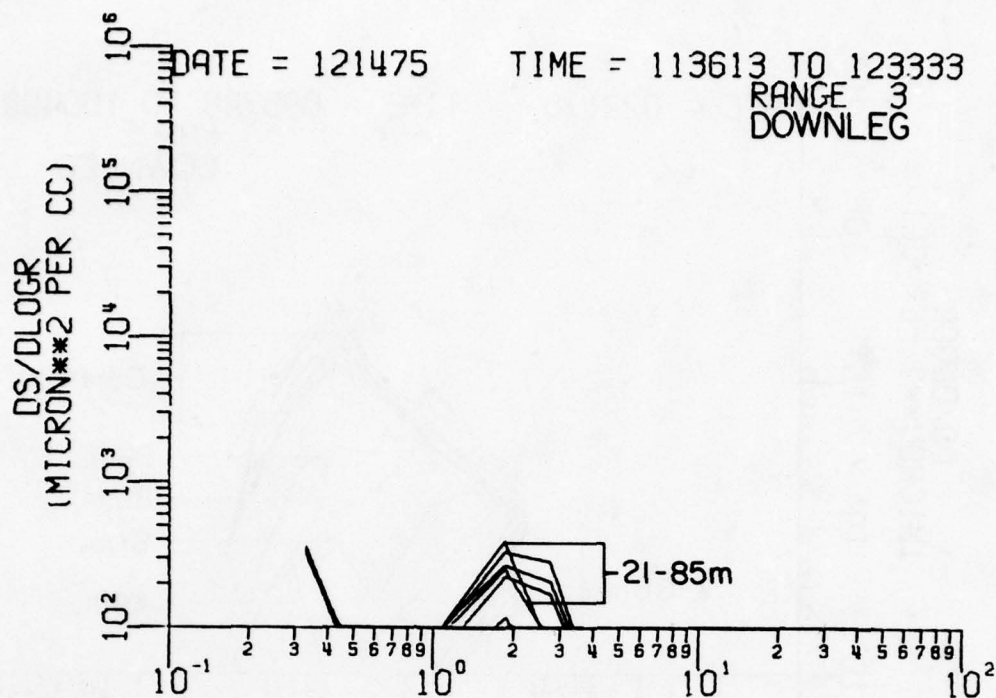


Figure A2-a. Same as Fig. A1 except for particle area (cross-sectional) distribution.

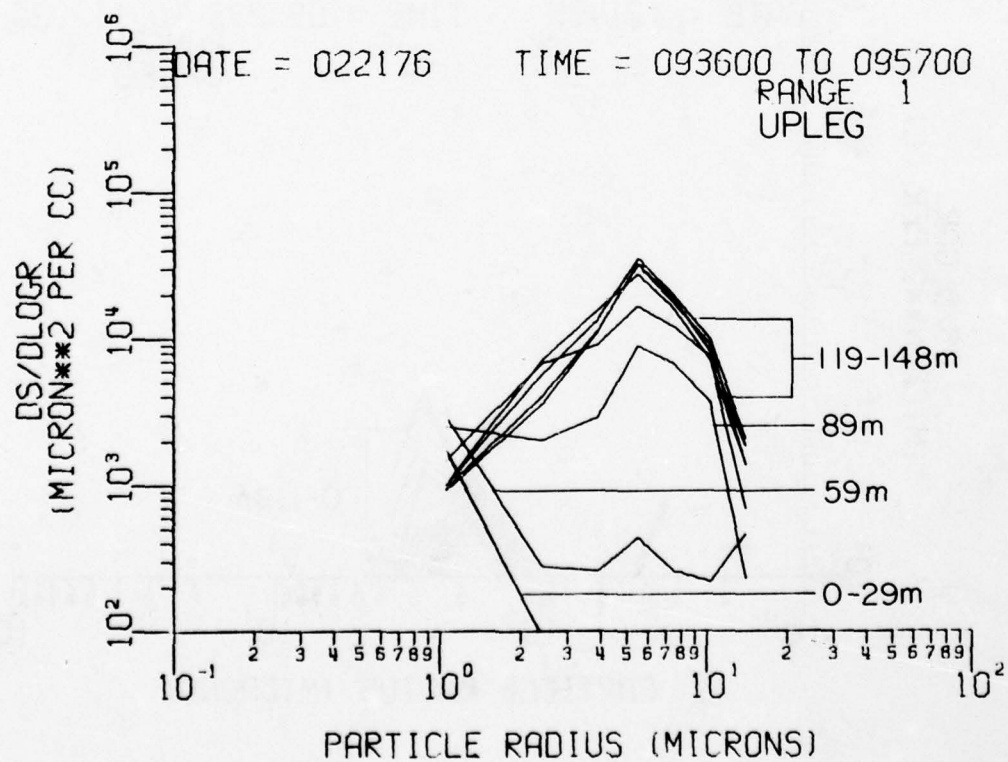
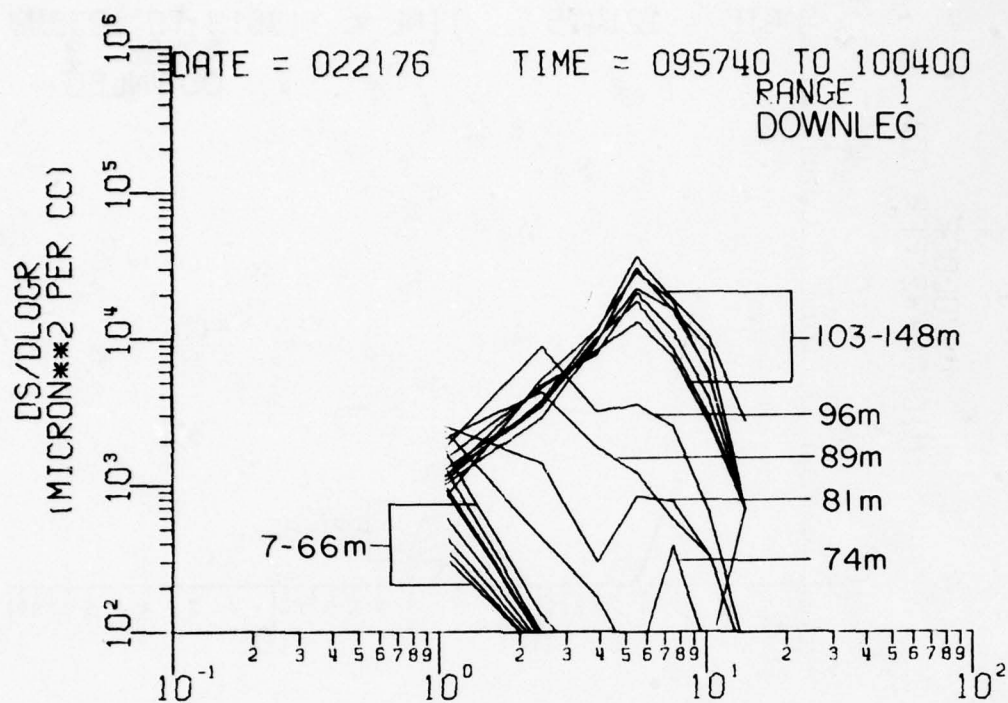


Figure A2-b

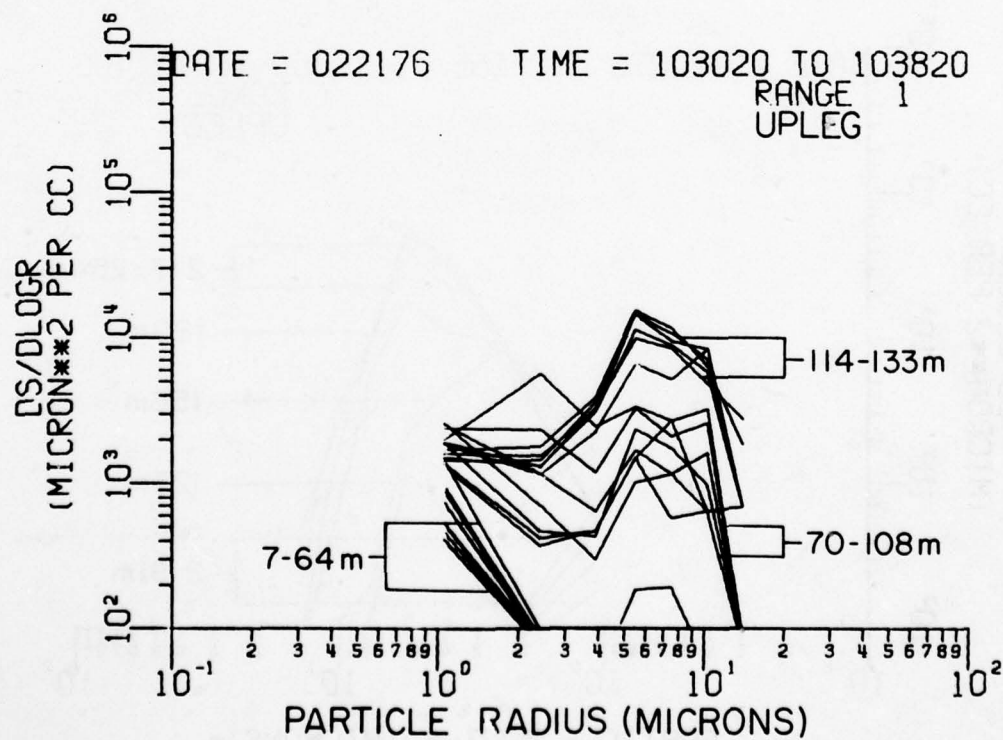
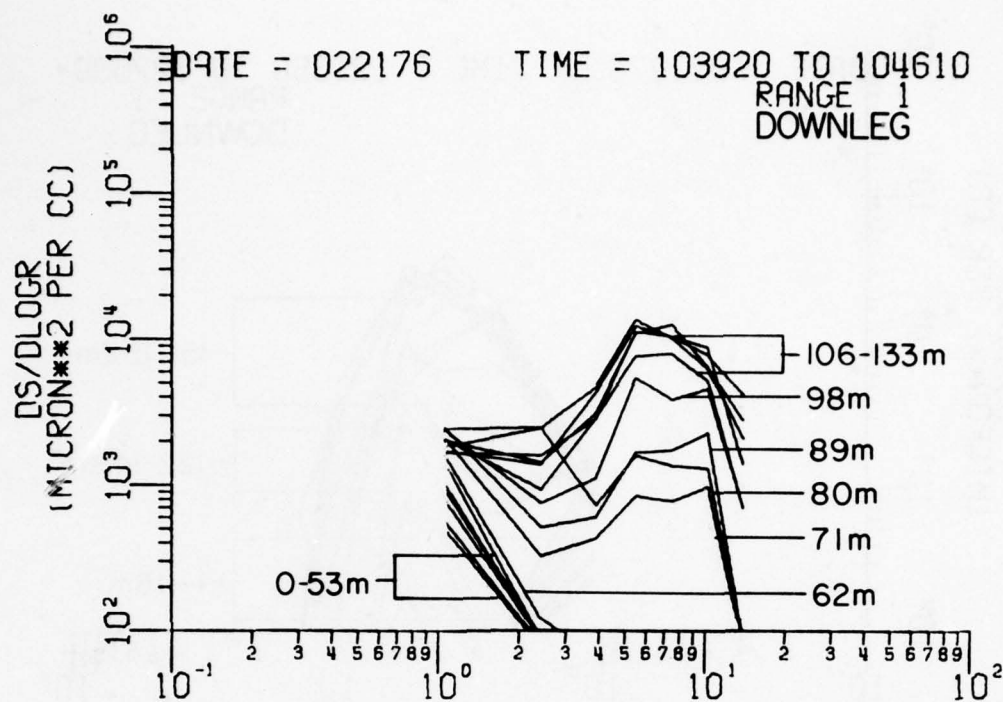


Figure A2-c

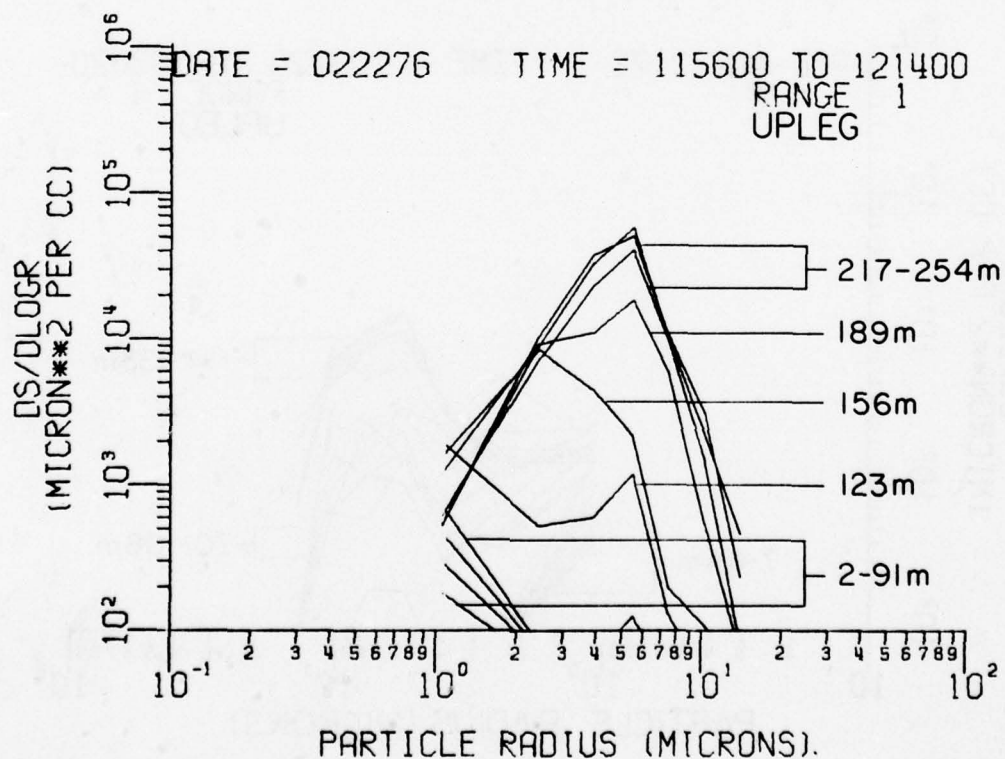
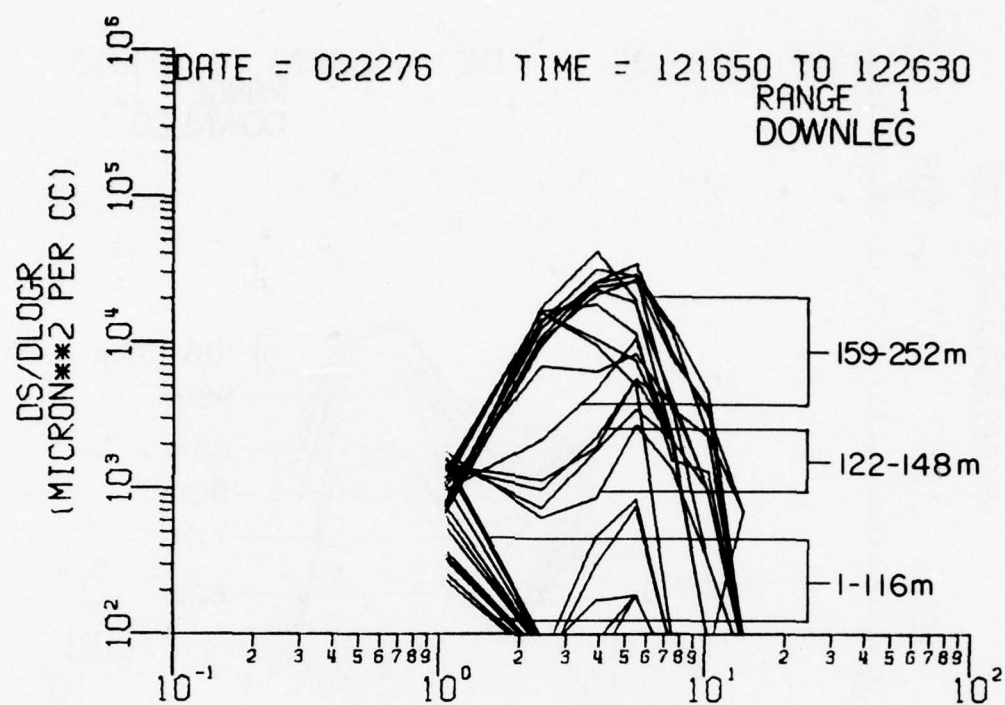


Figure A2-d

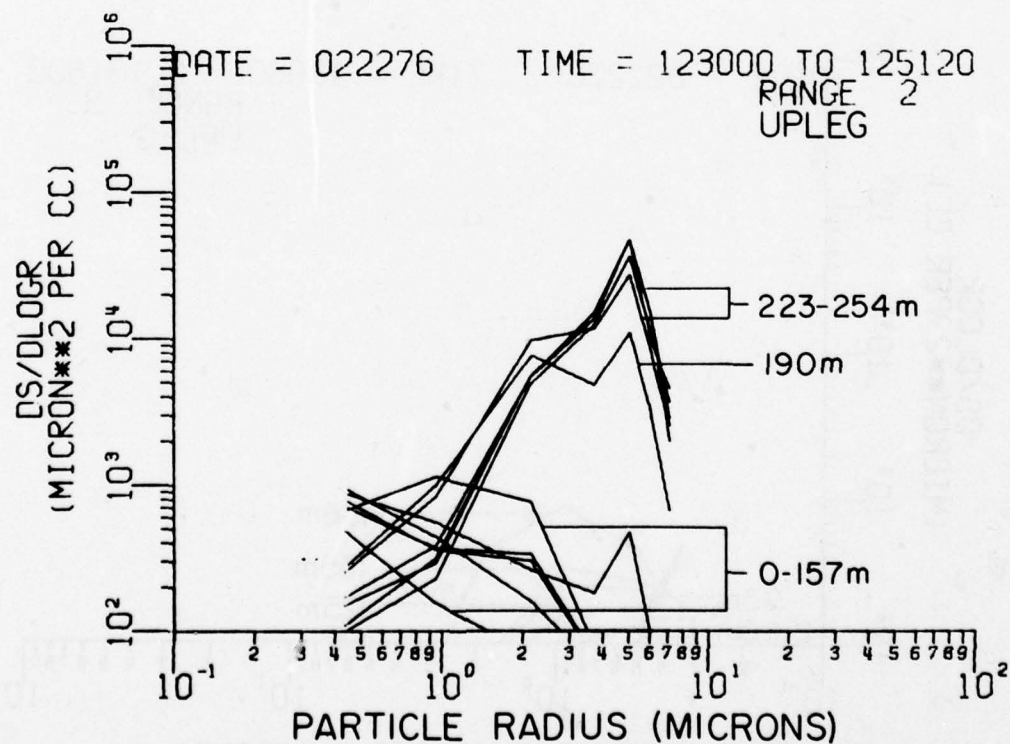
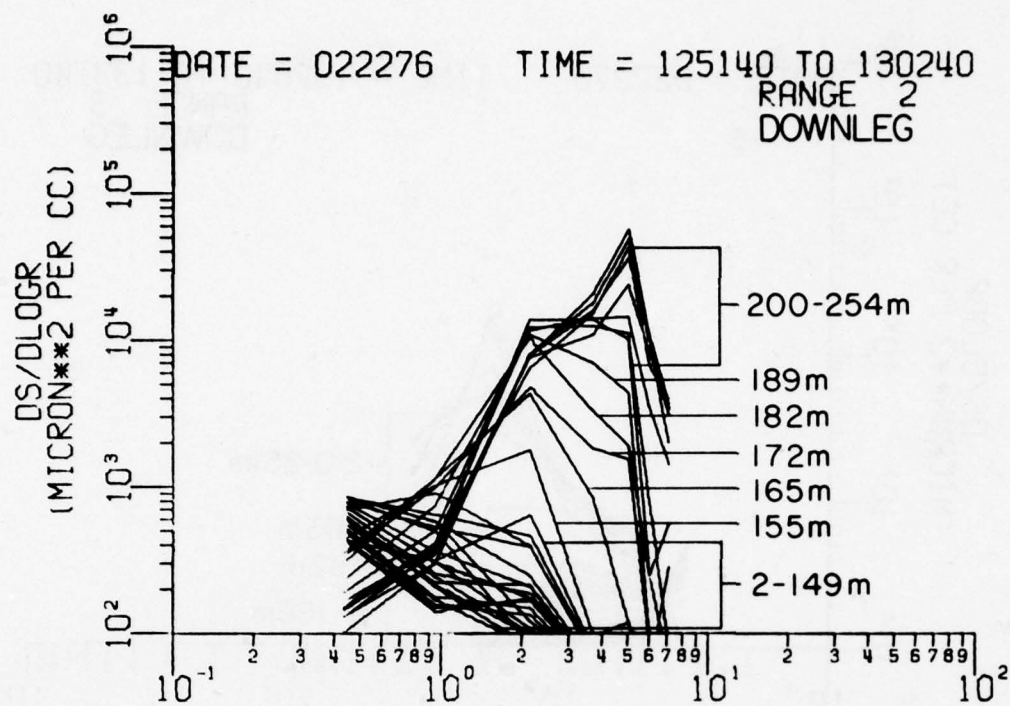


Figure A2-e

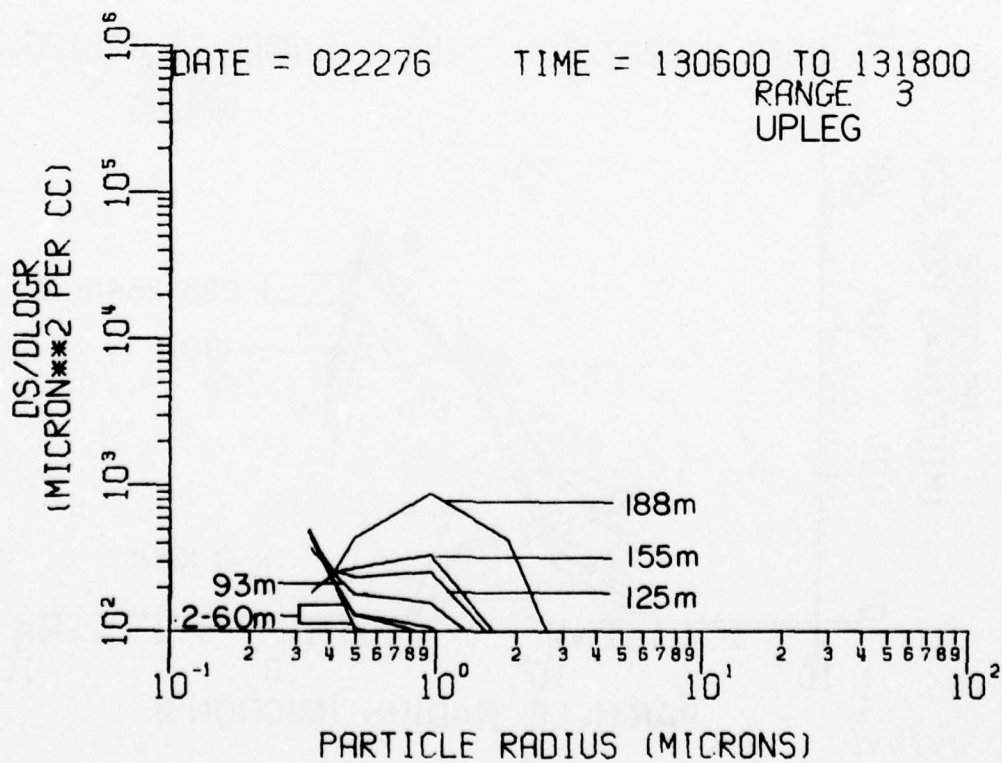
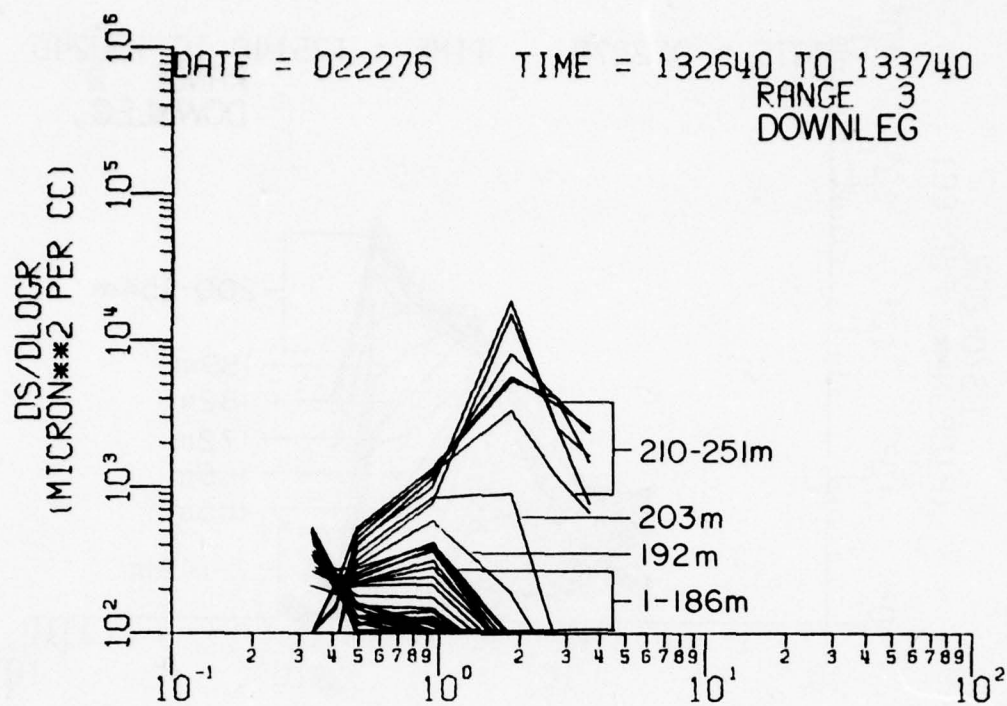


Figure A2-f

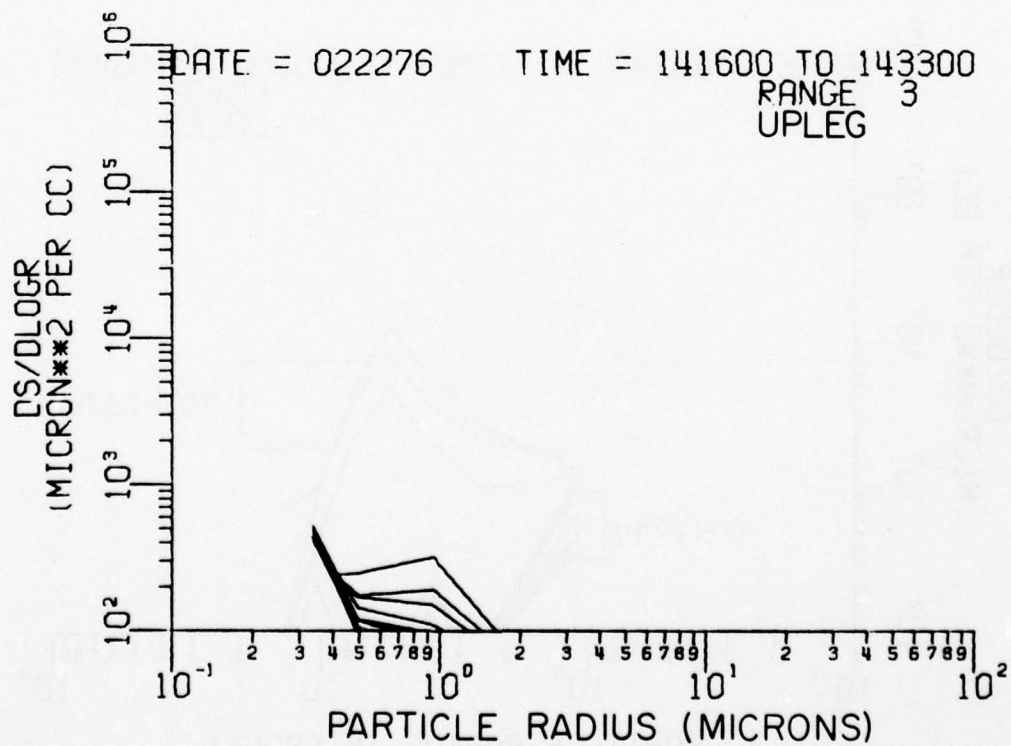
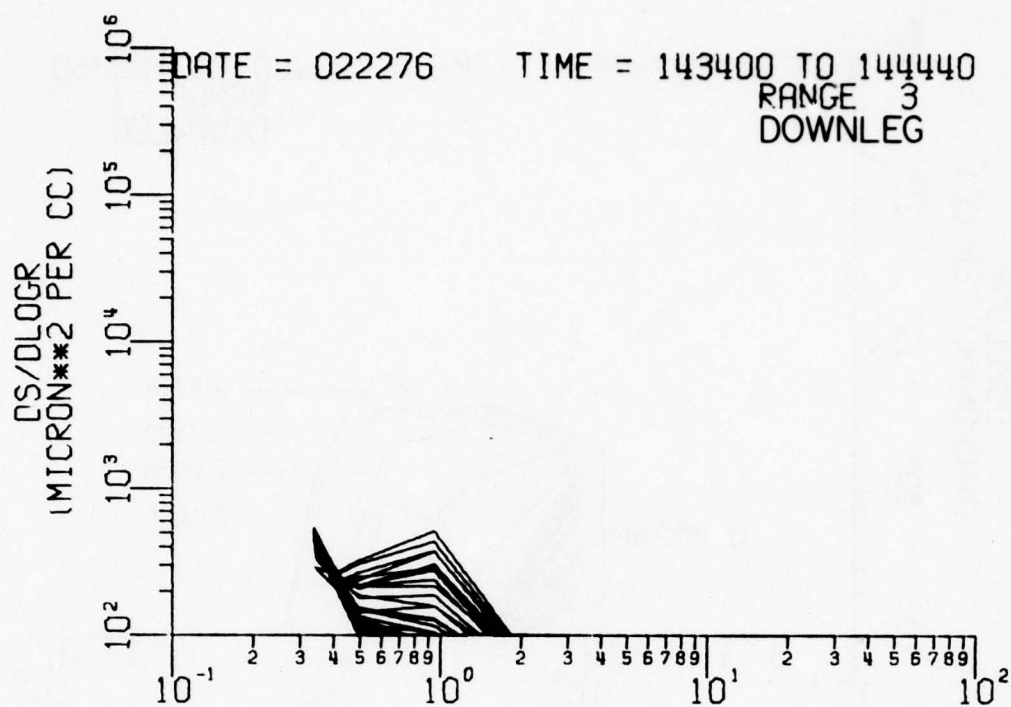


Figure A2-g

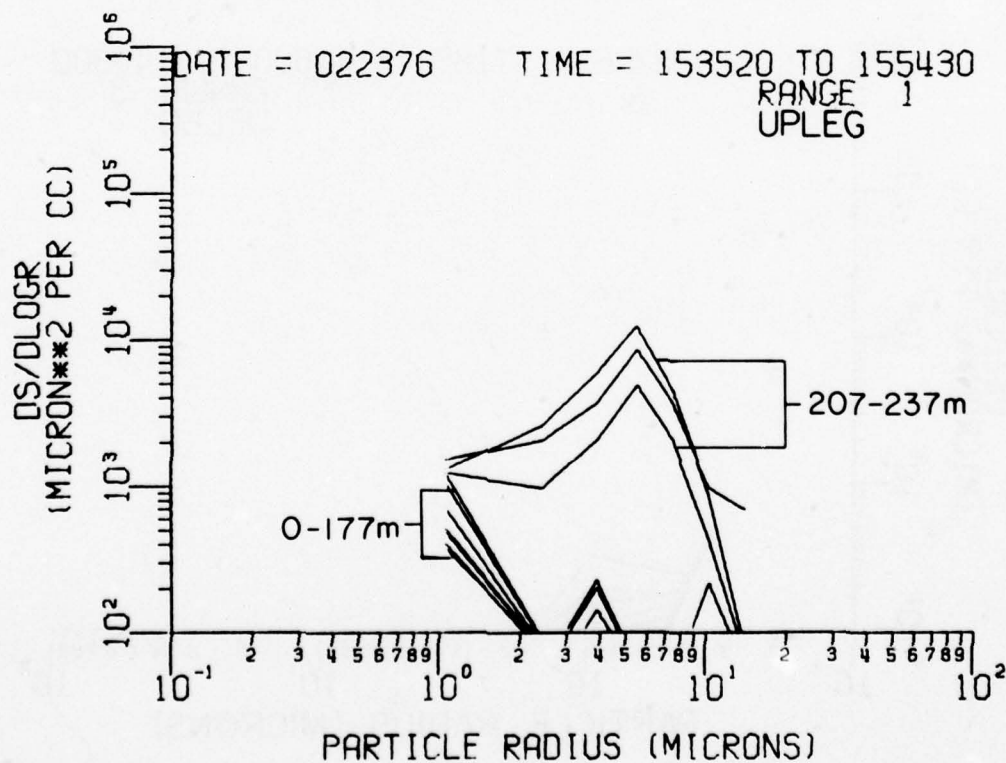
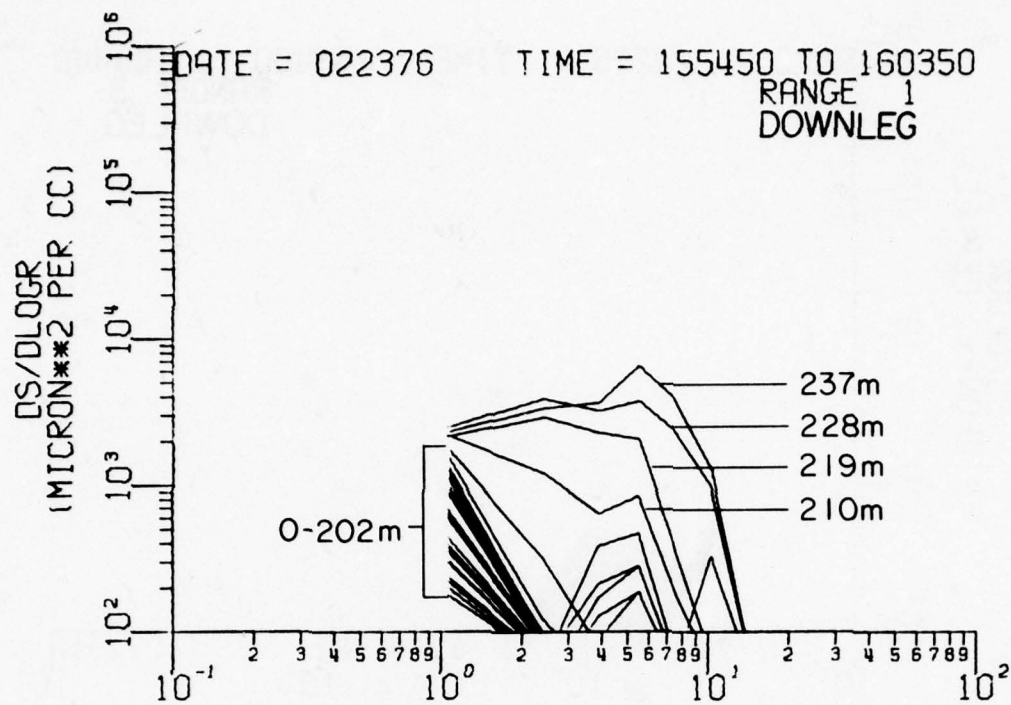


Figure A2-h

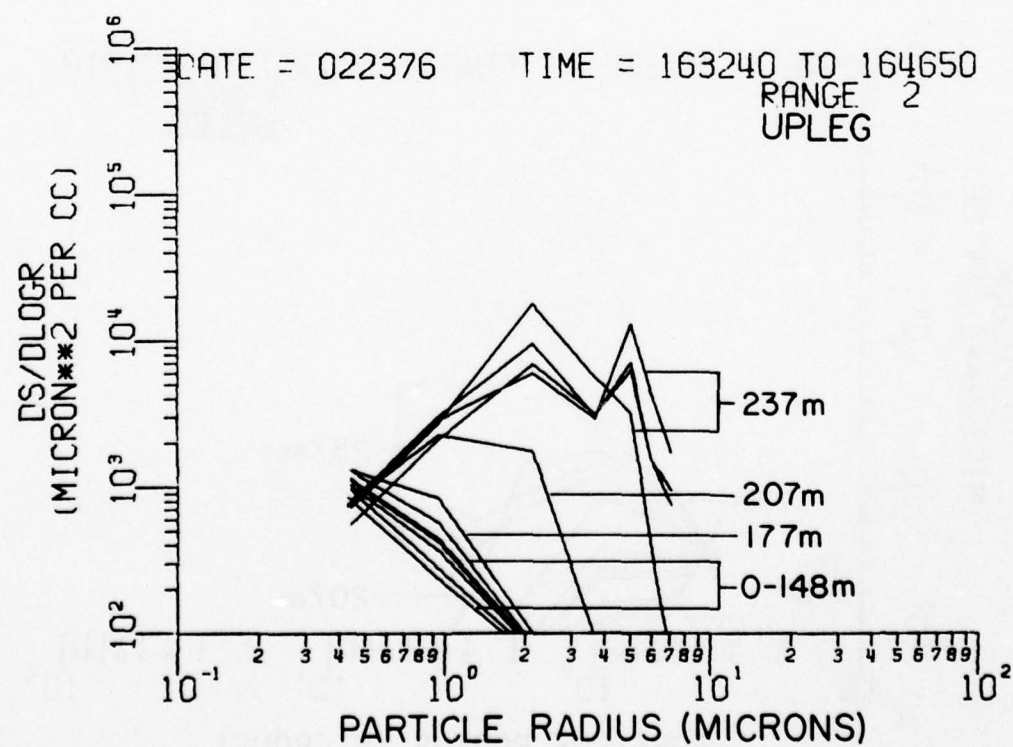
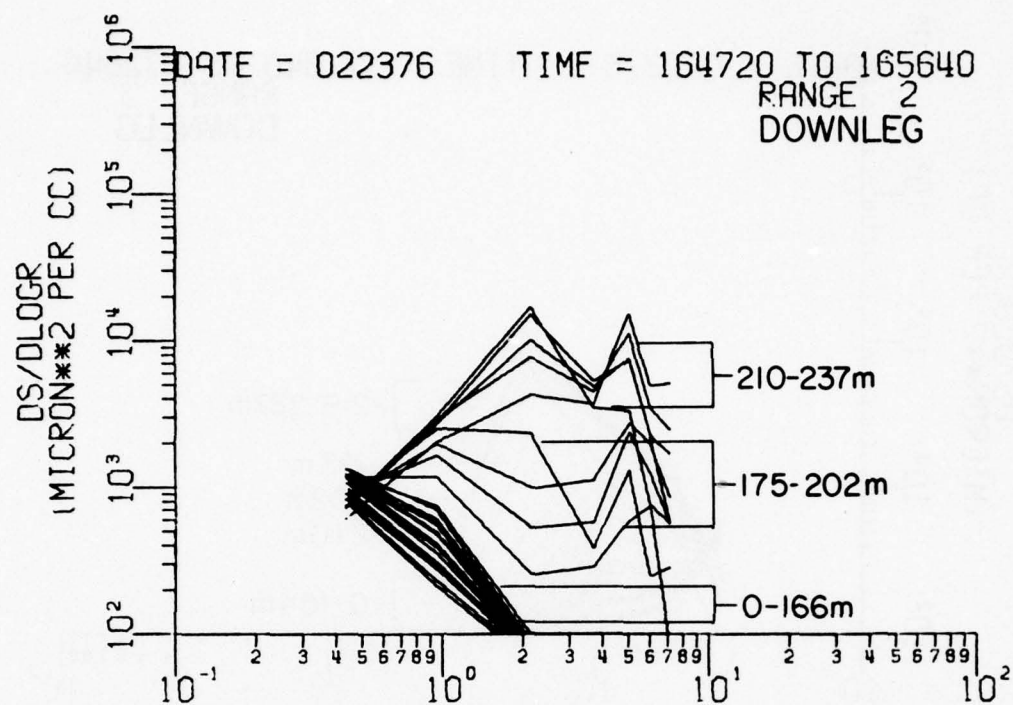


Figure A2-i

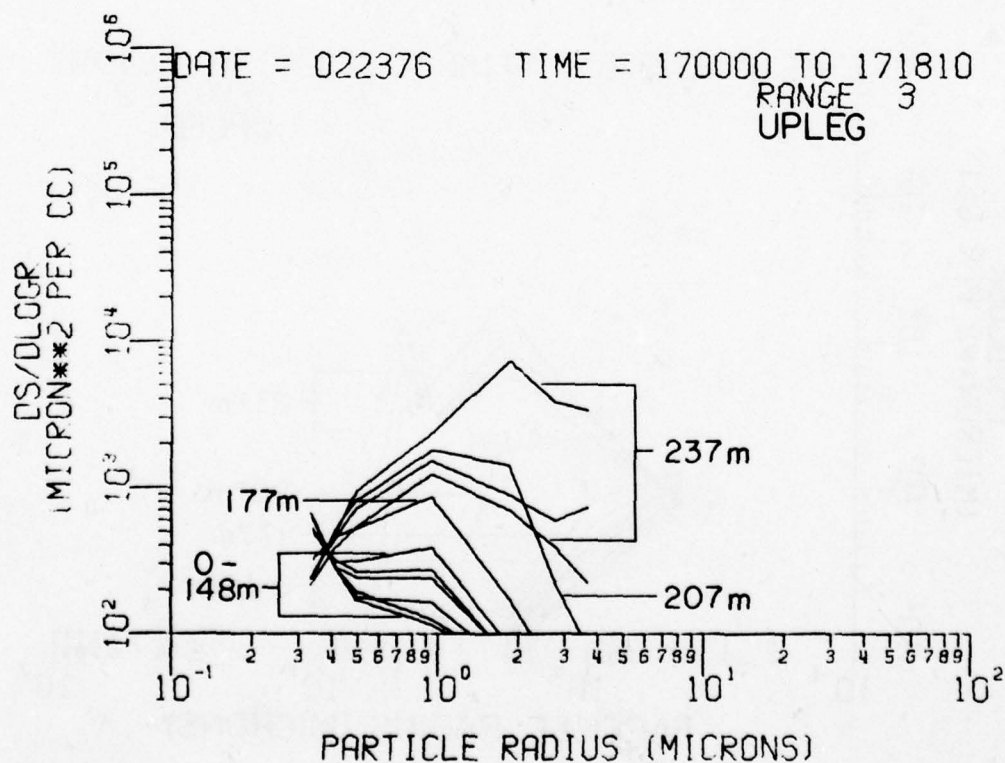
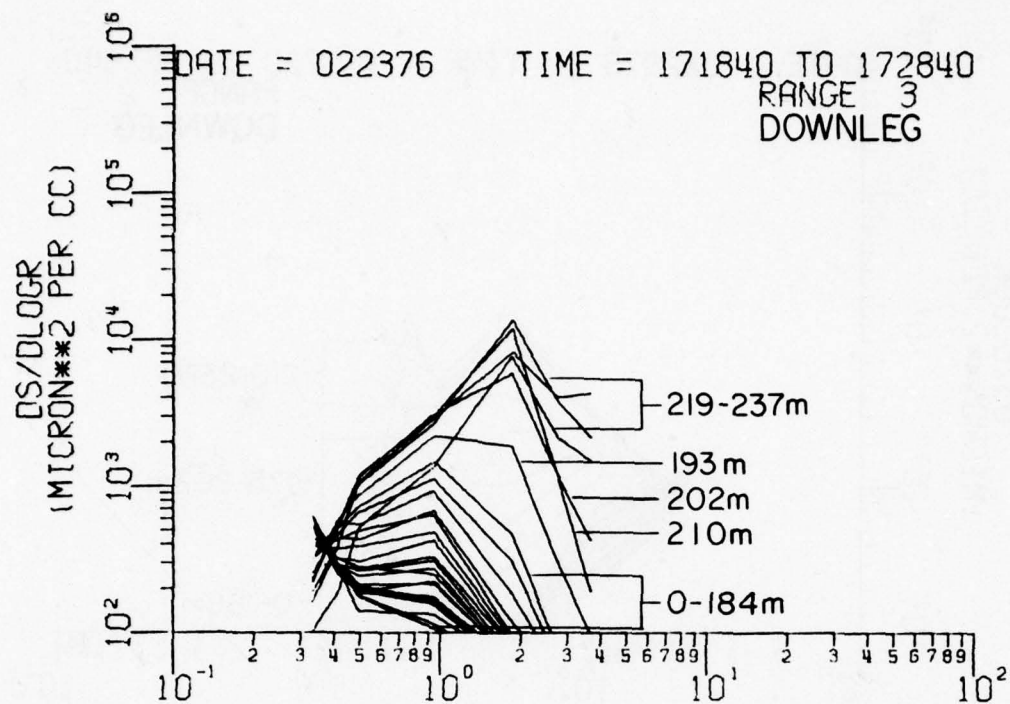


Figure A2-j

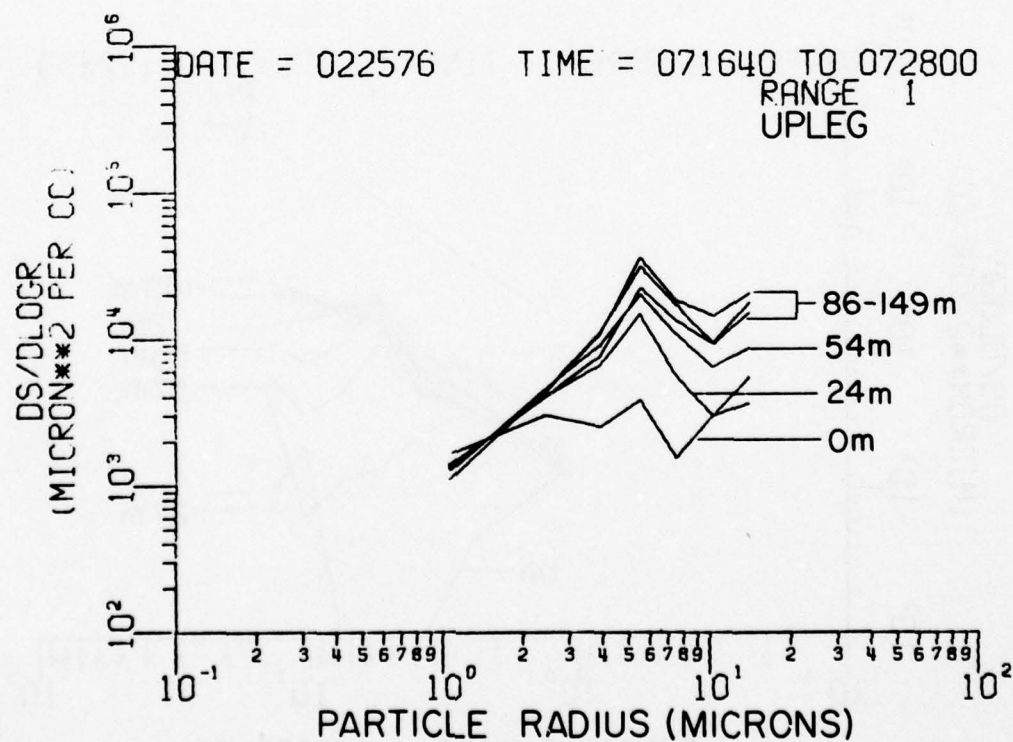
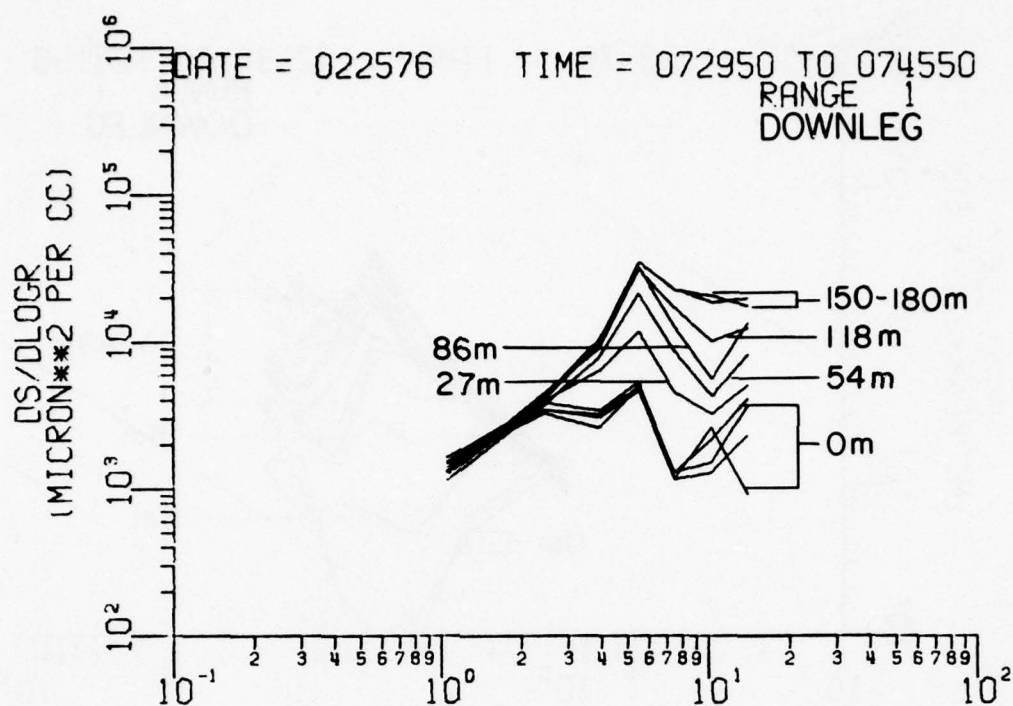


Figure A2-k

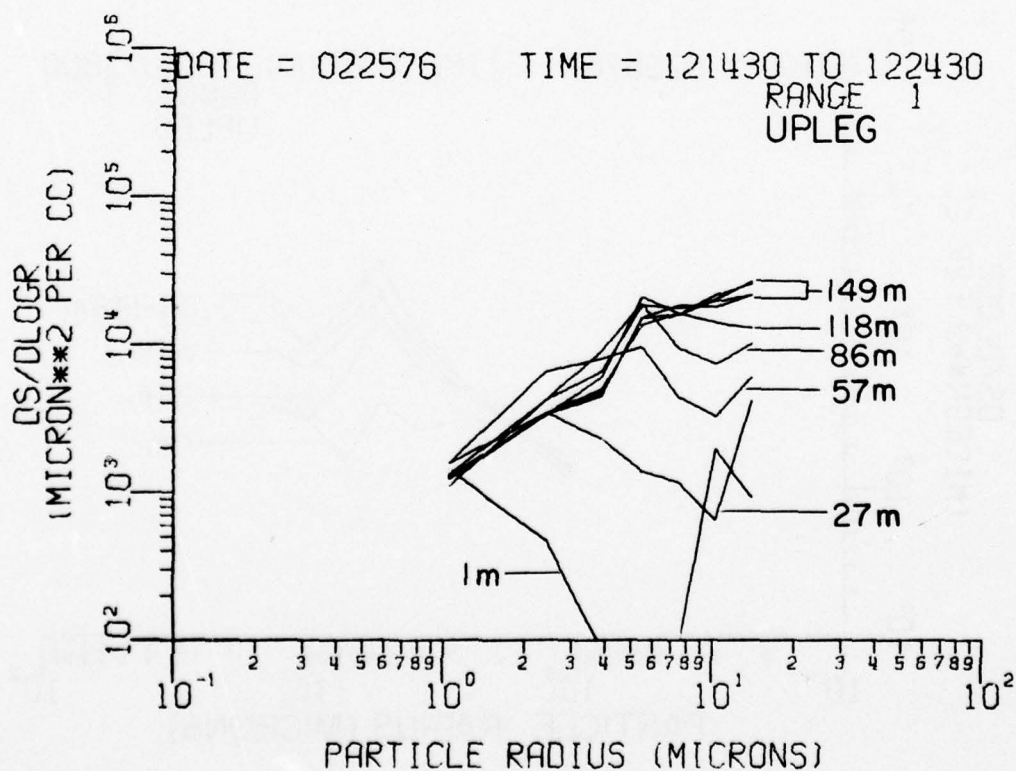
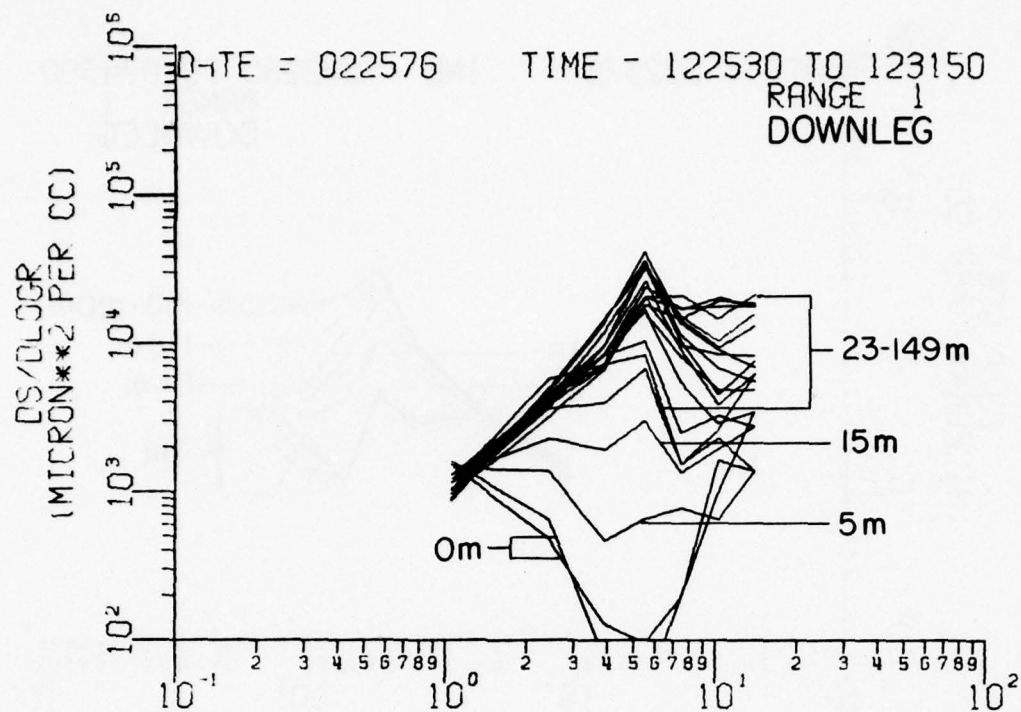


Figure A2-1

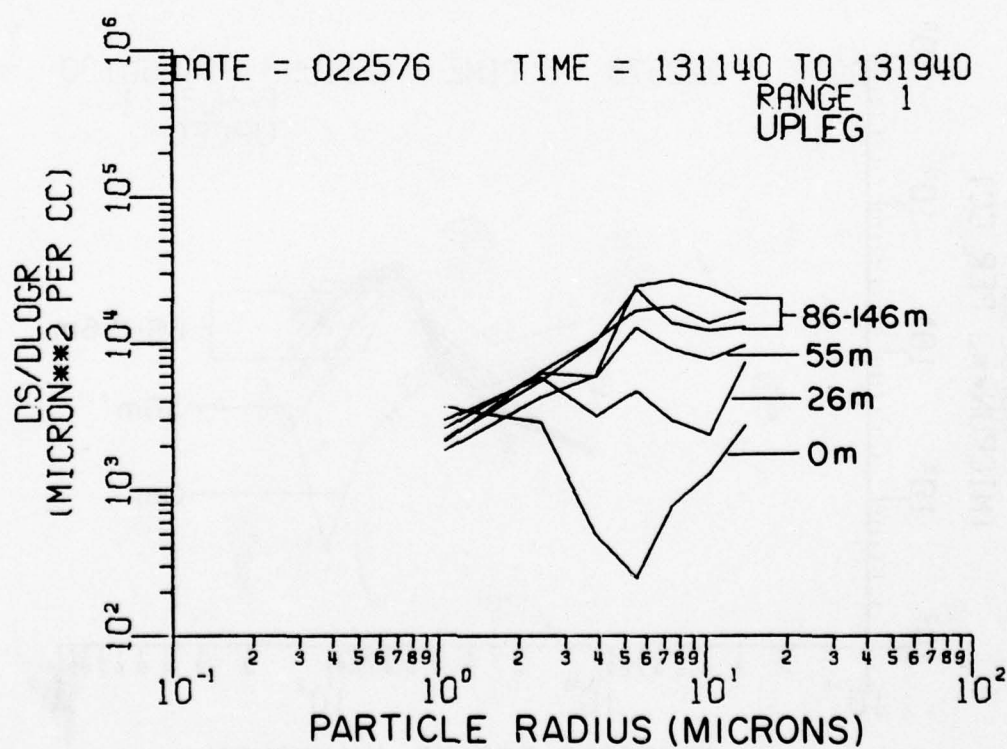
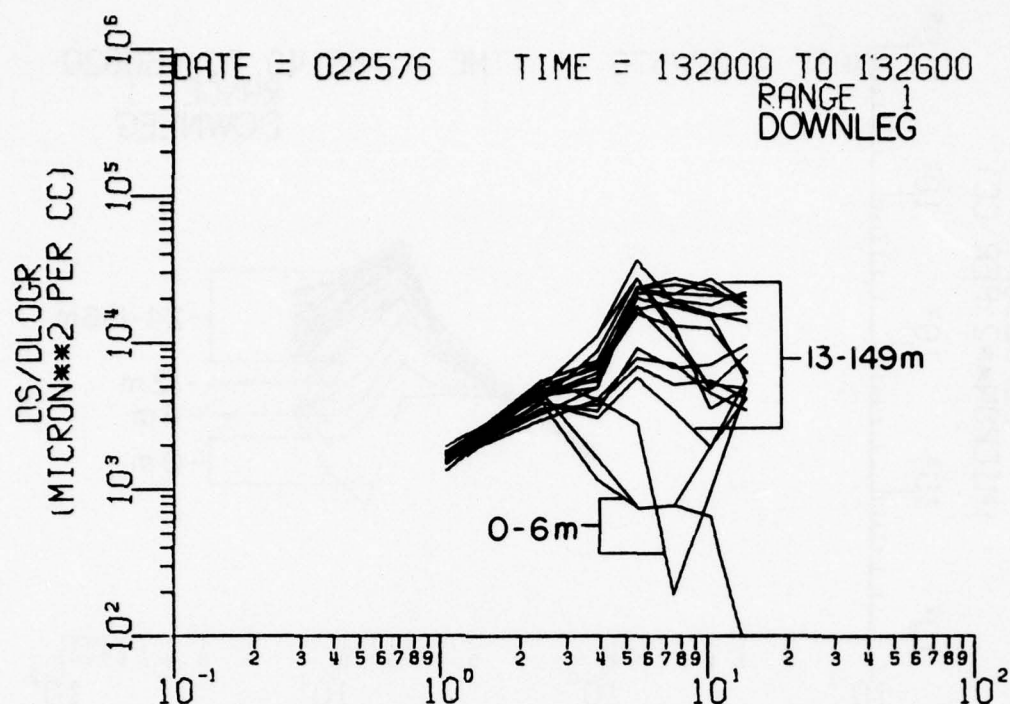


Figure A2-m

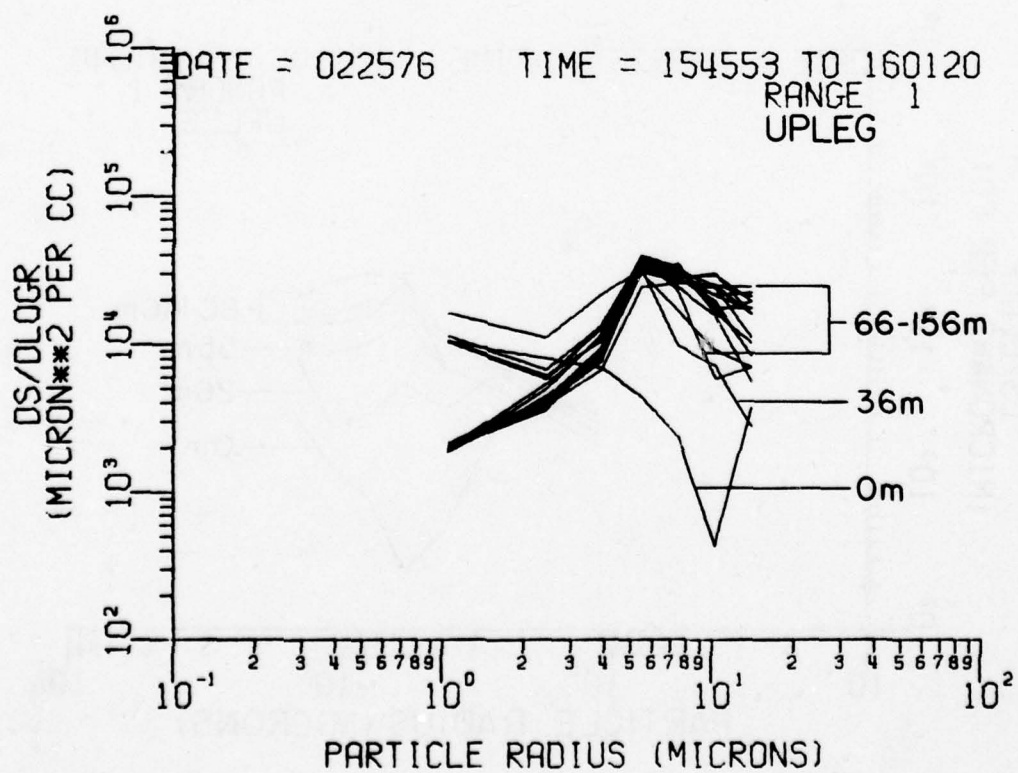
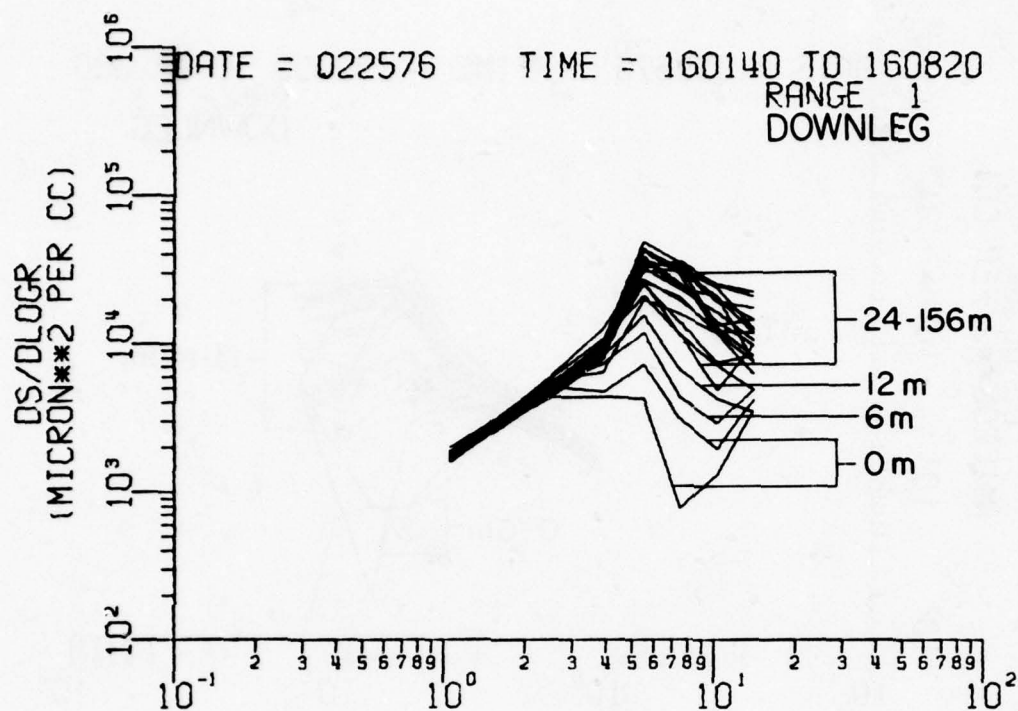


Figure A2-n

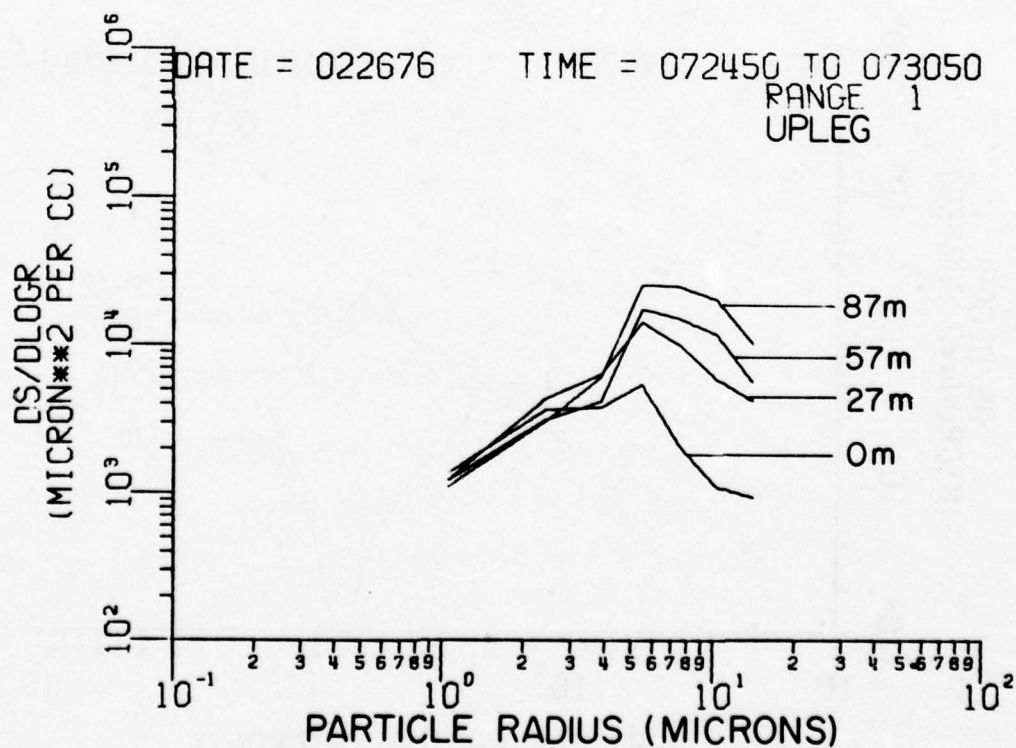
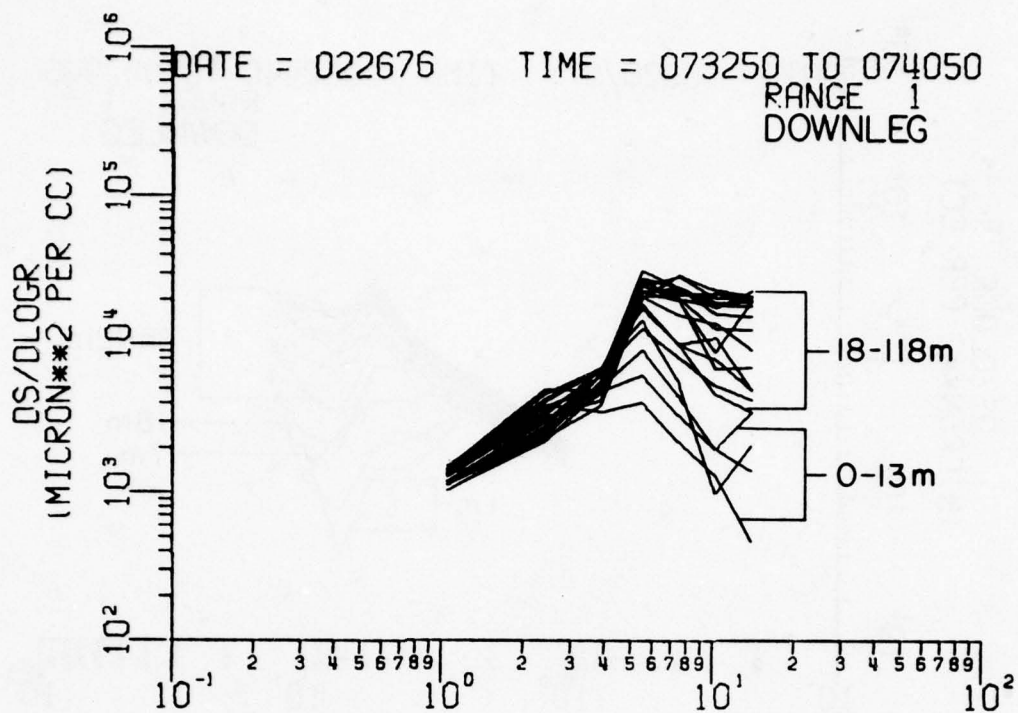


Figure A2-o

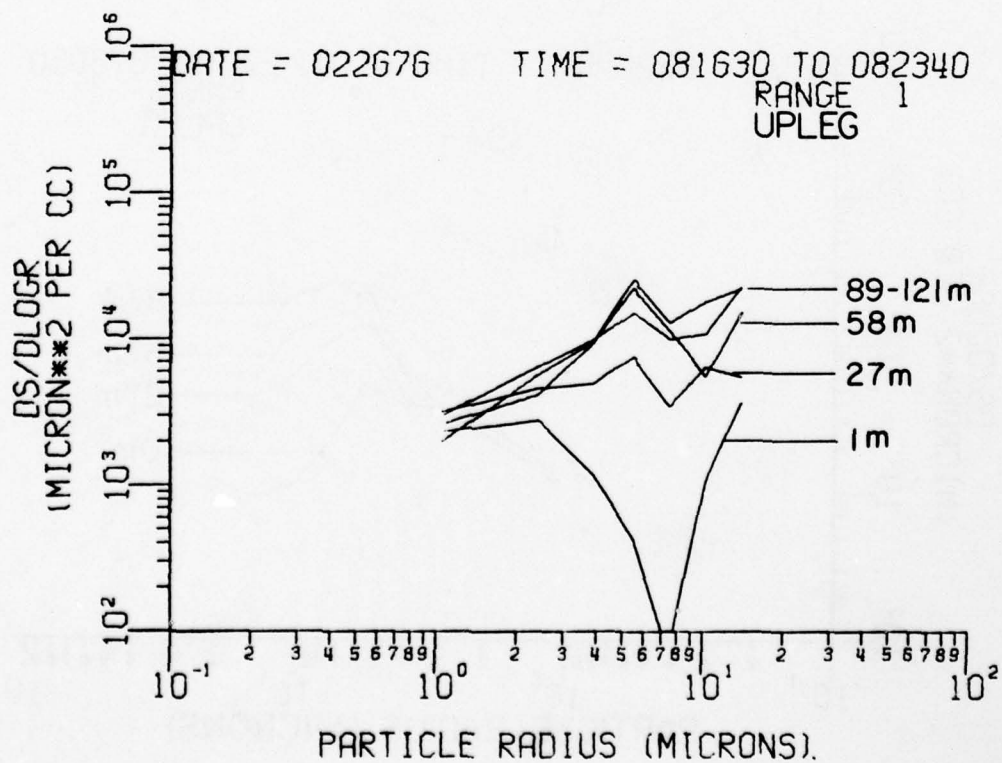
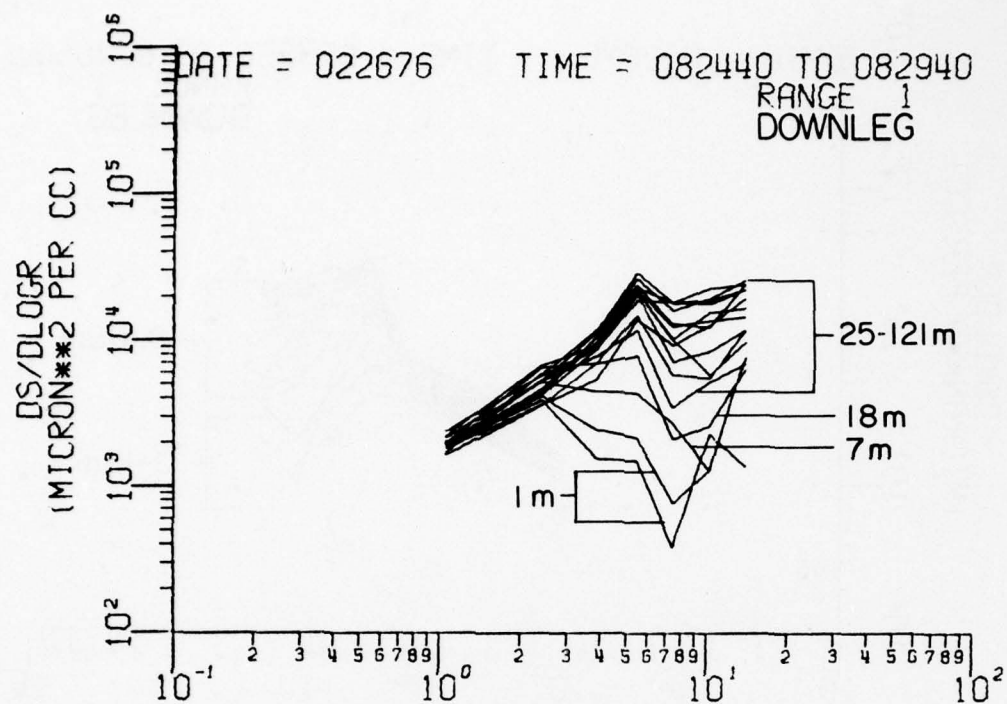


Figure A2-p

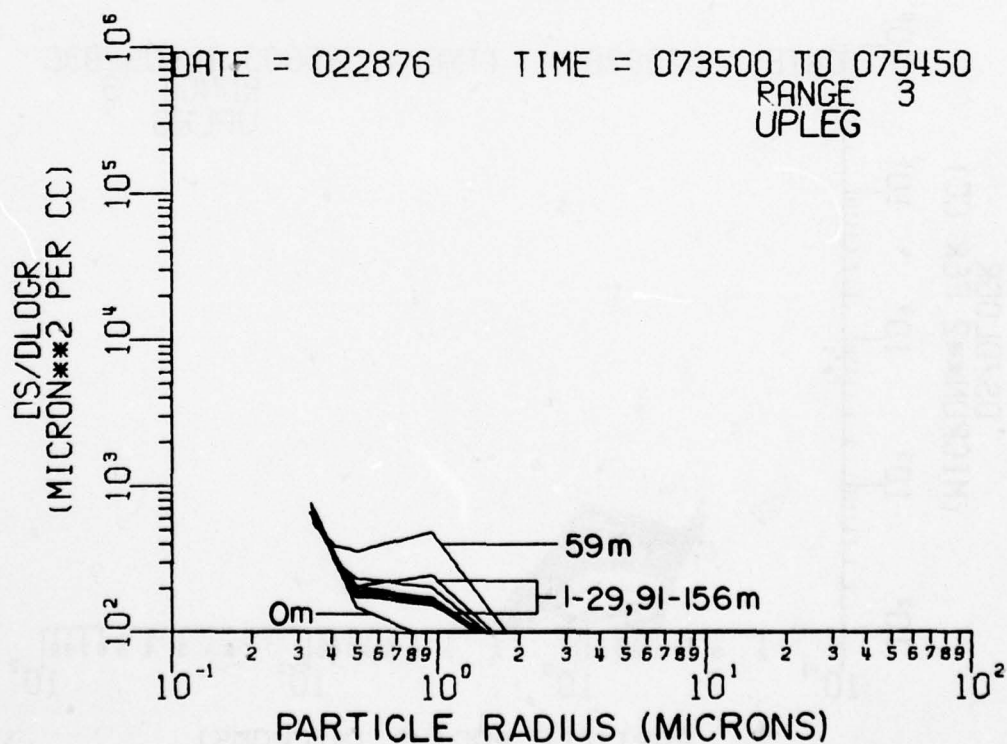
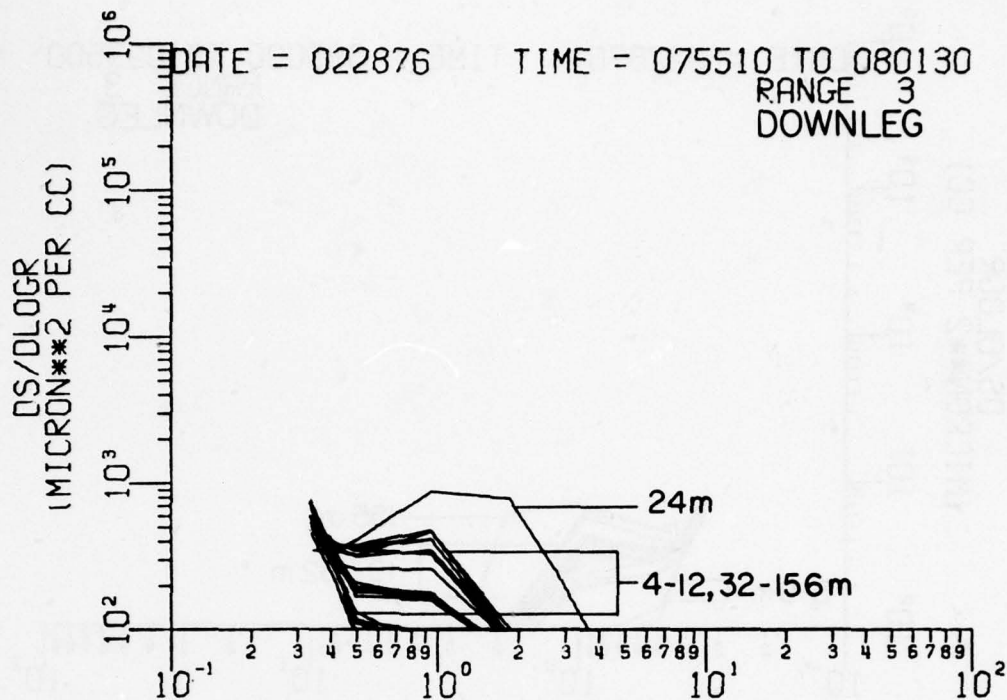


Figure A2-q

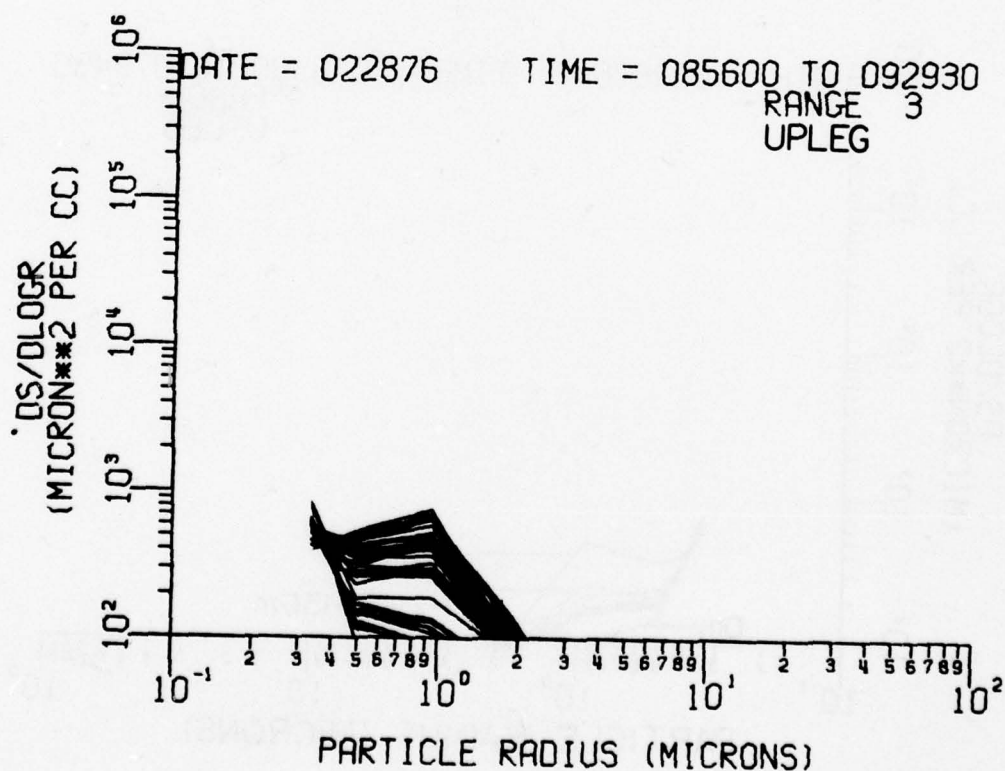
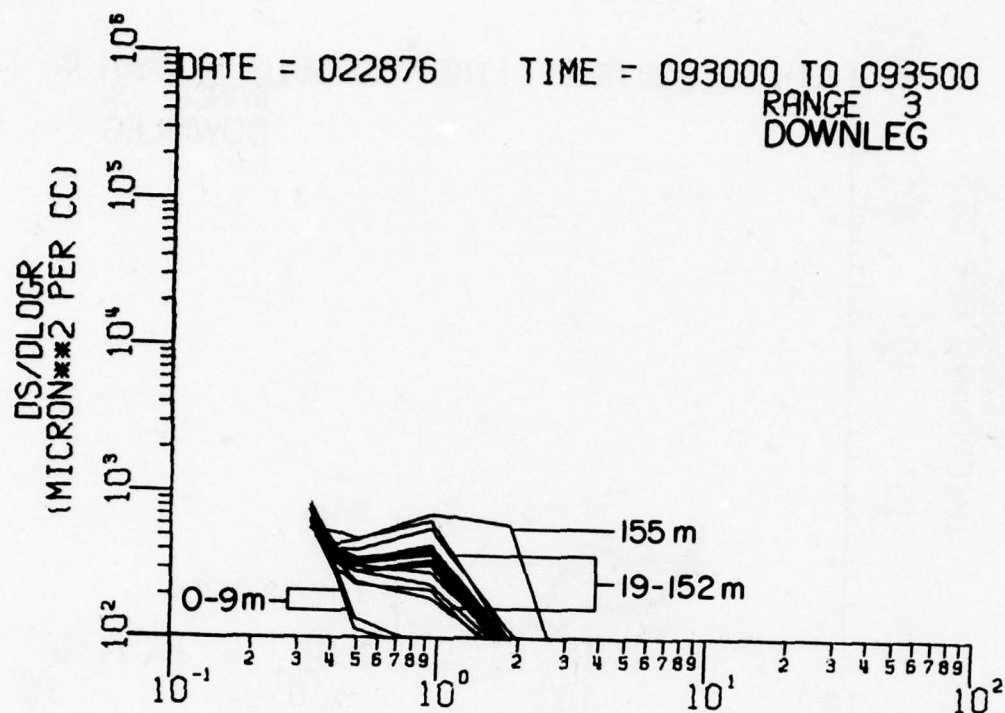


Figure A2-r

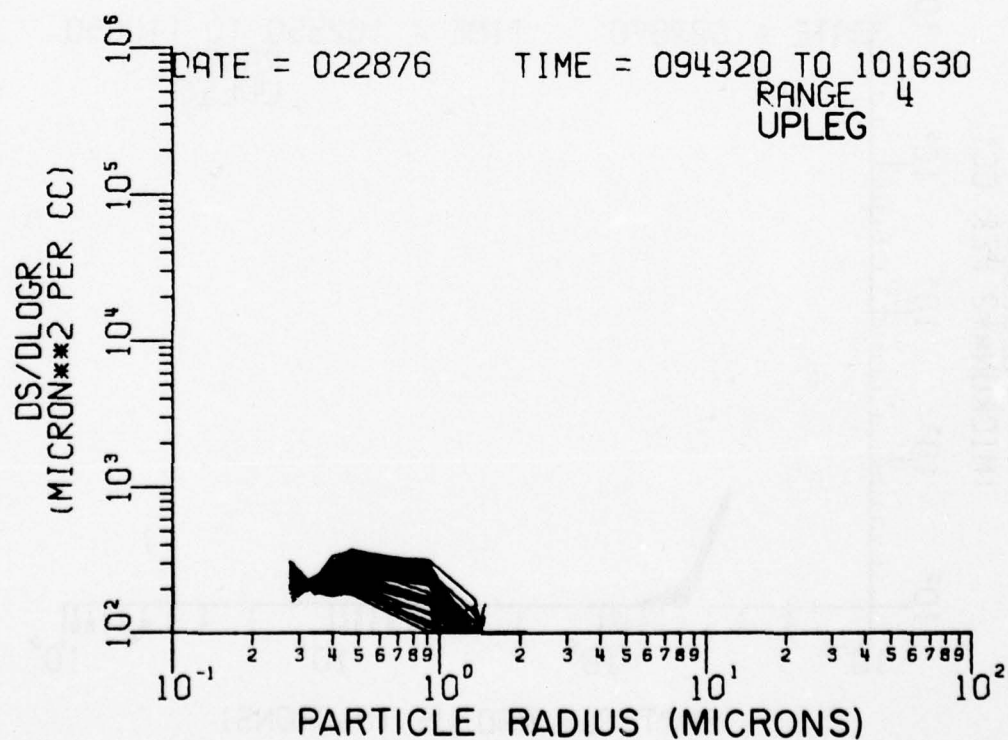
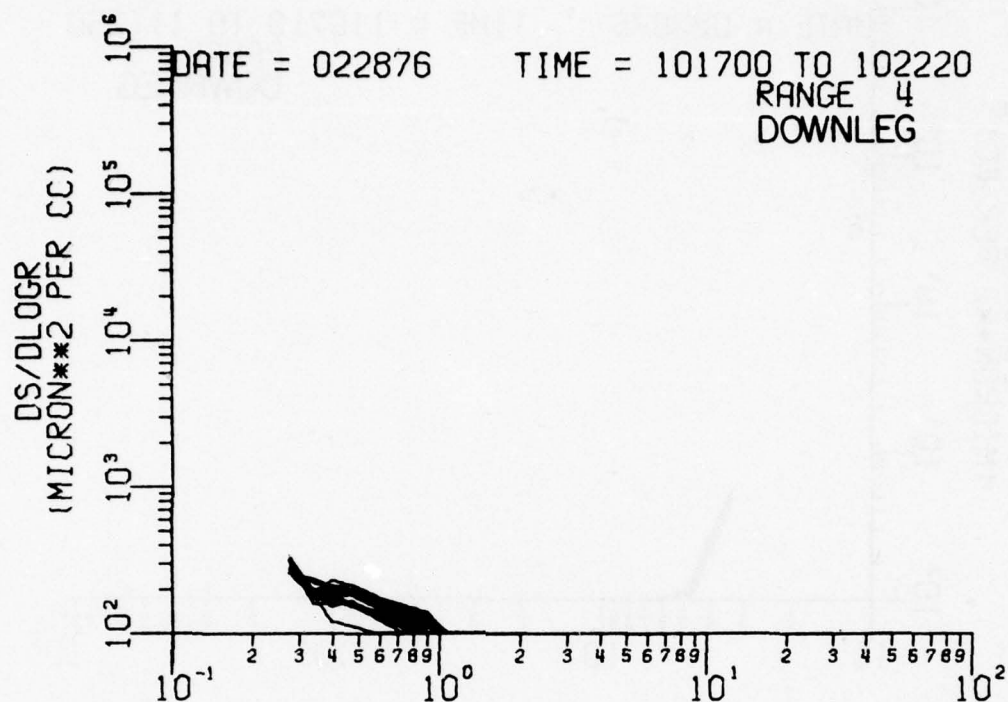


Figure A2-s

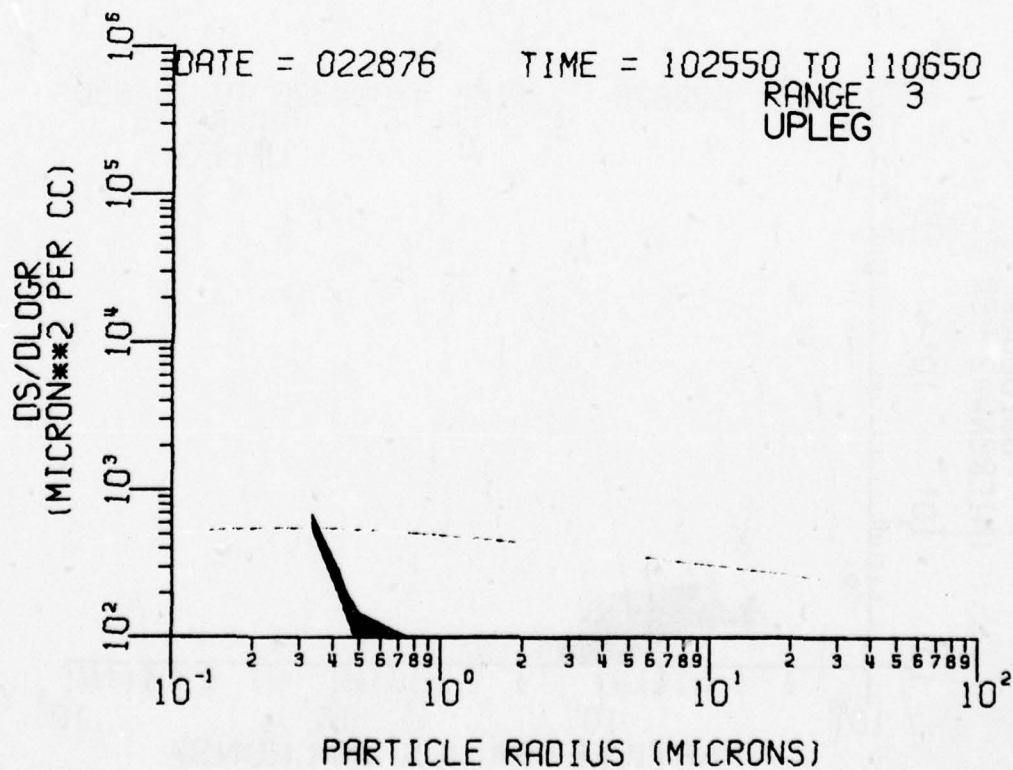
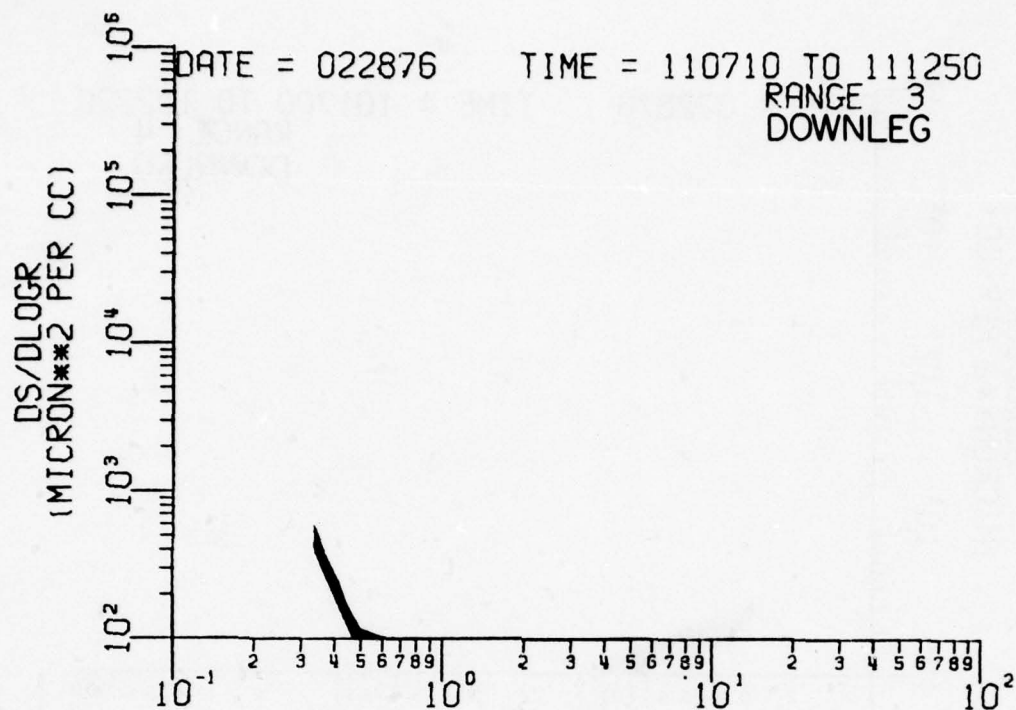


Figure A2-t

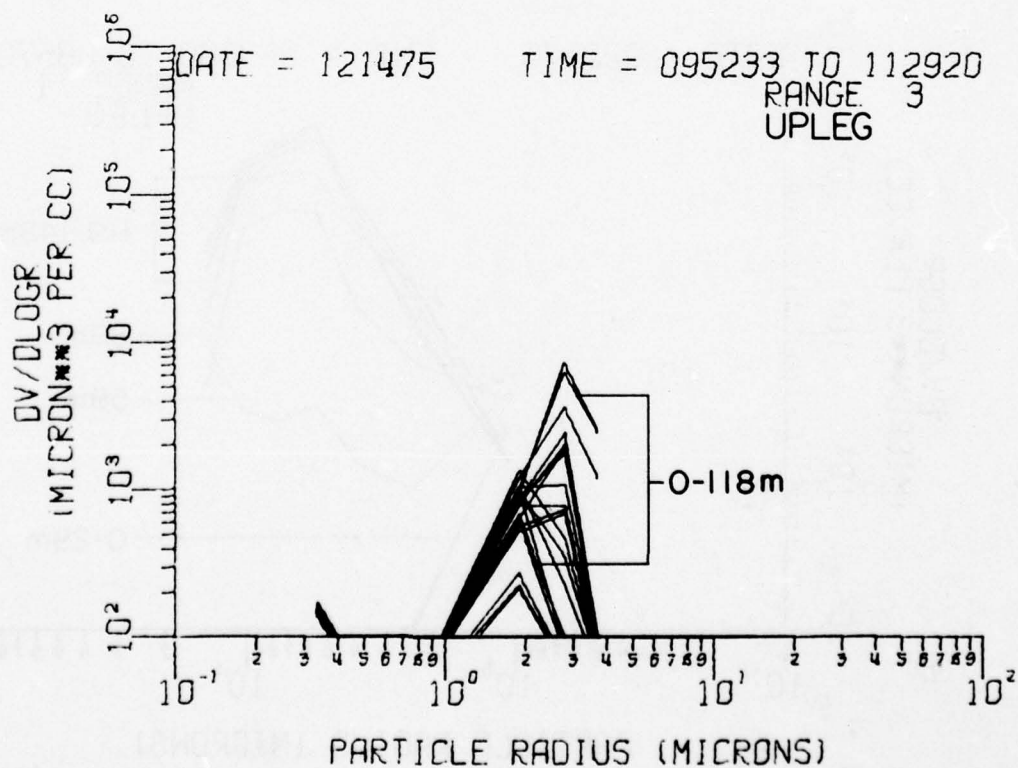
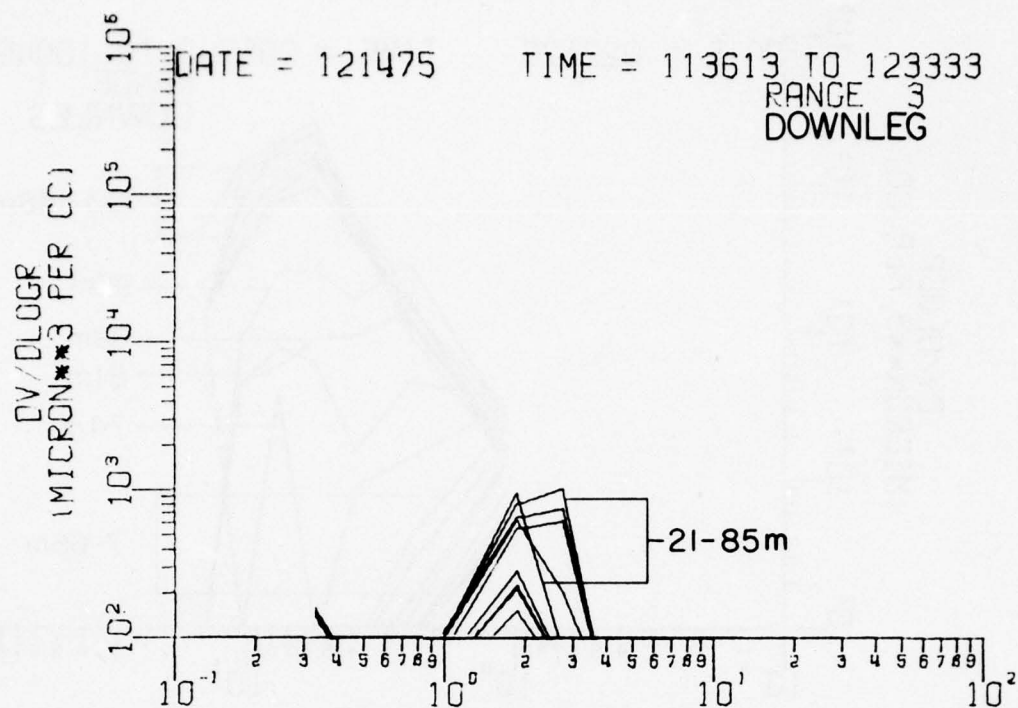


Figure A3-a. Same as Fig. A1 except for particle volume distribution.

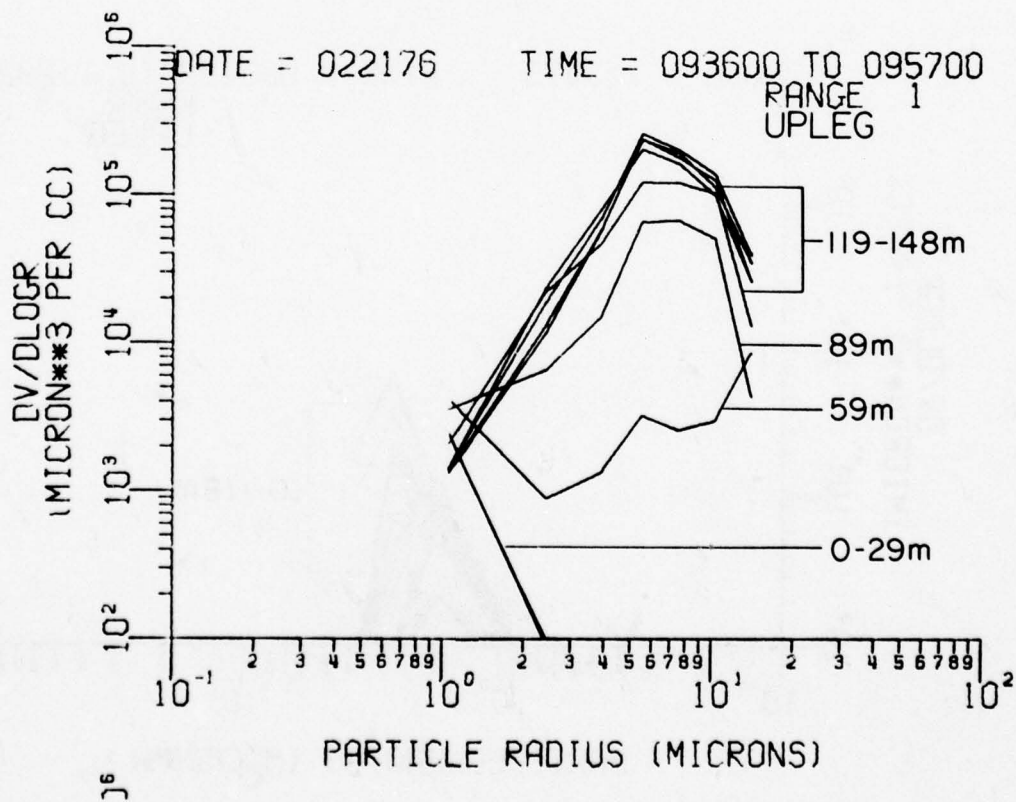
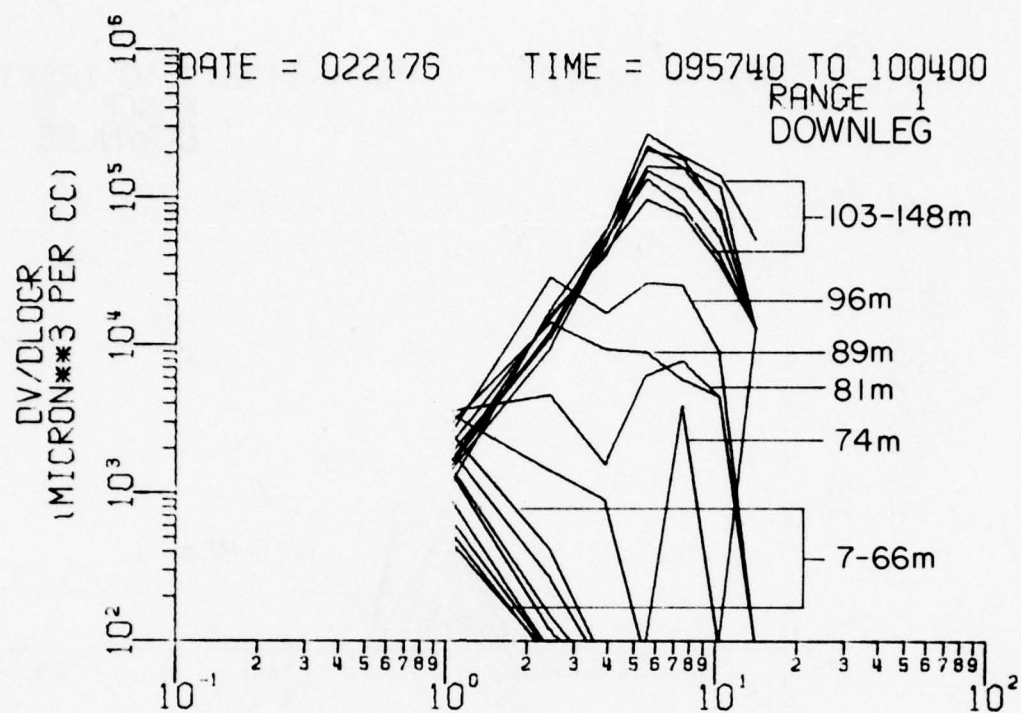


Figure A3-b

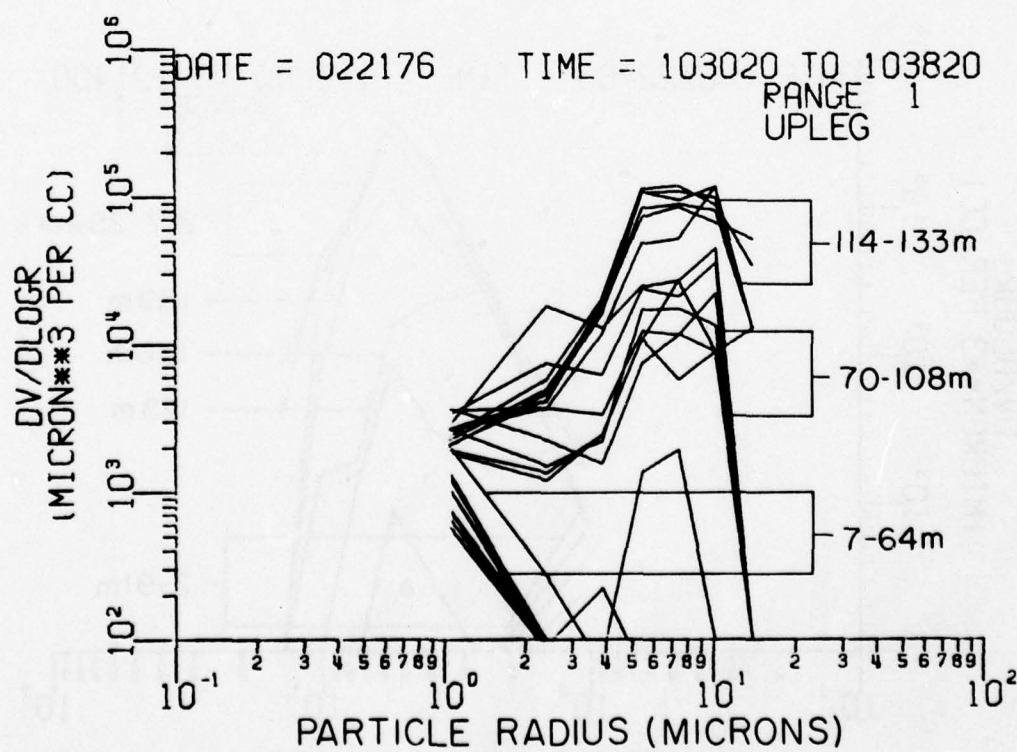
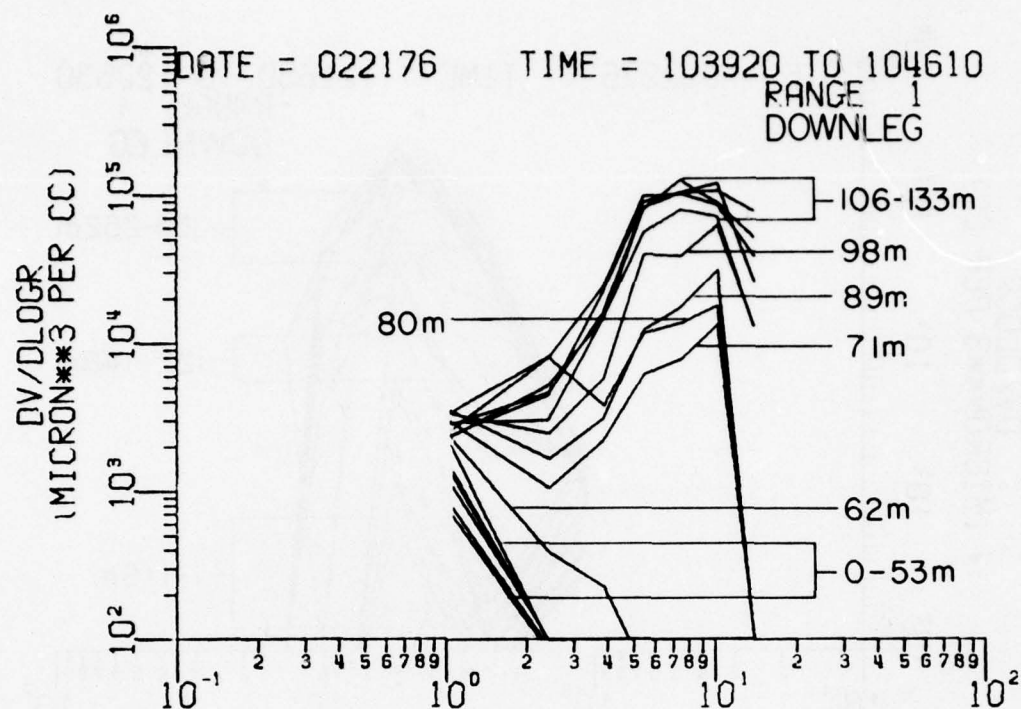


Figure A3-c

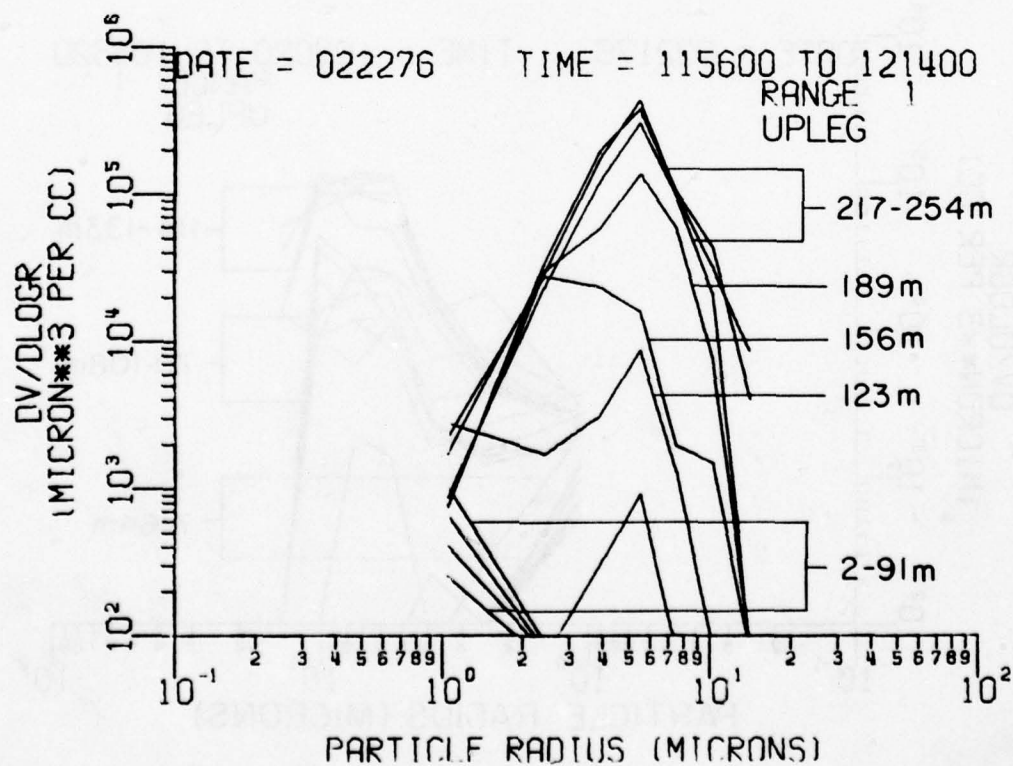
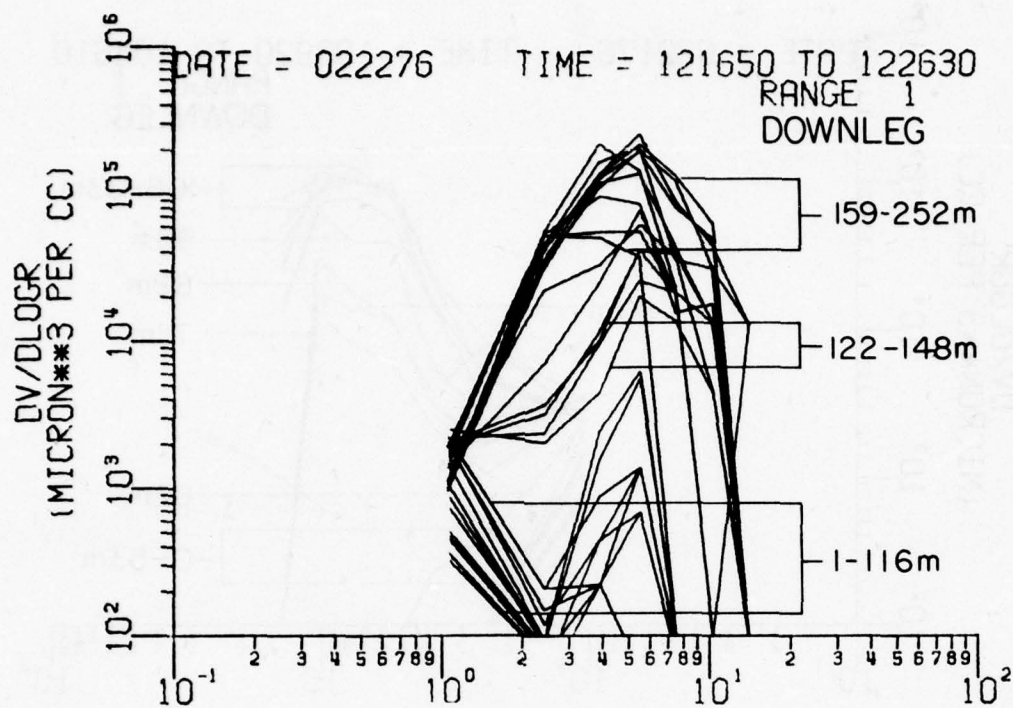


Figure A3-d

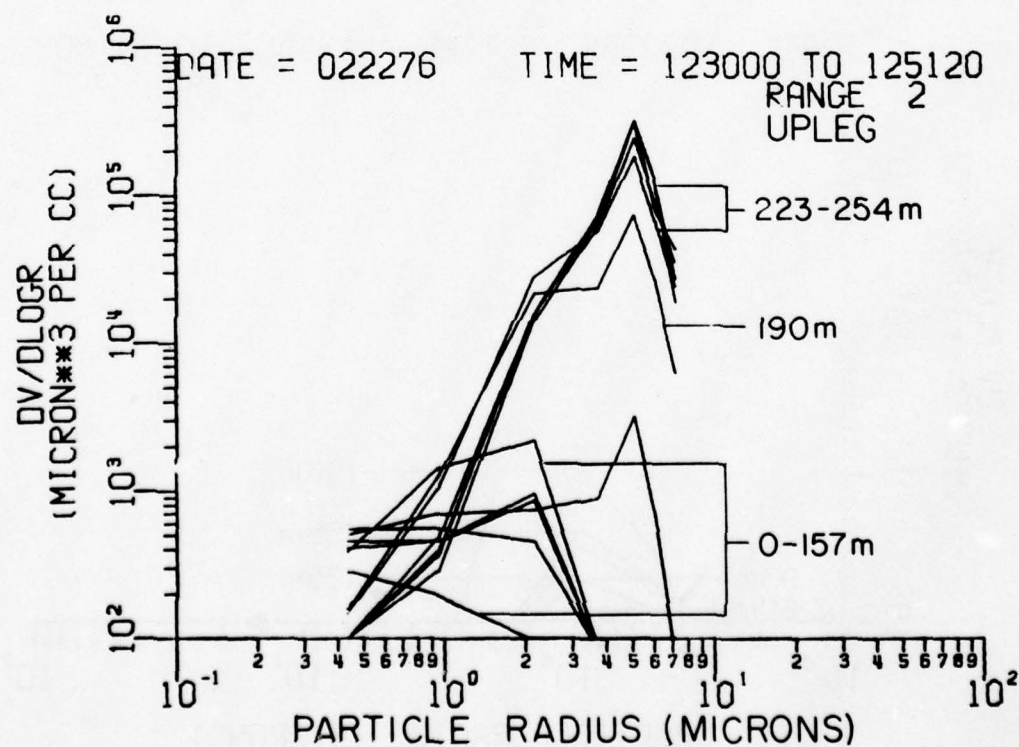
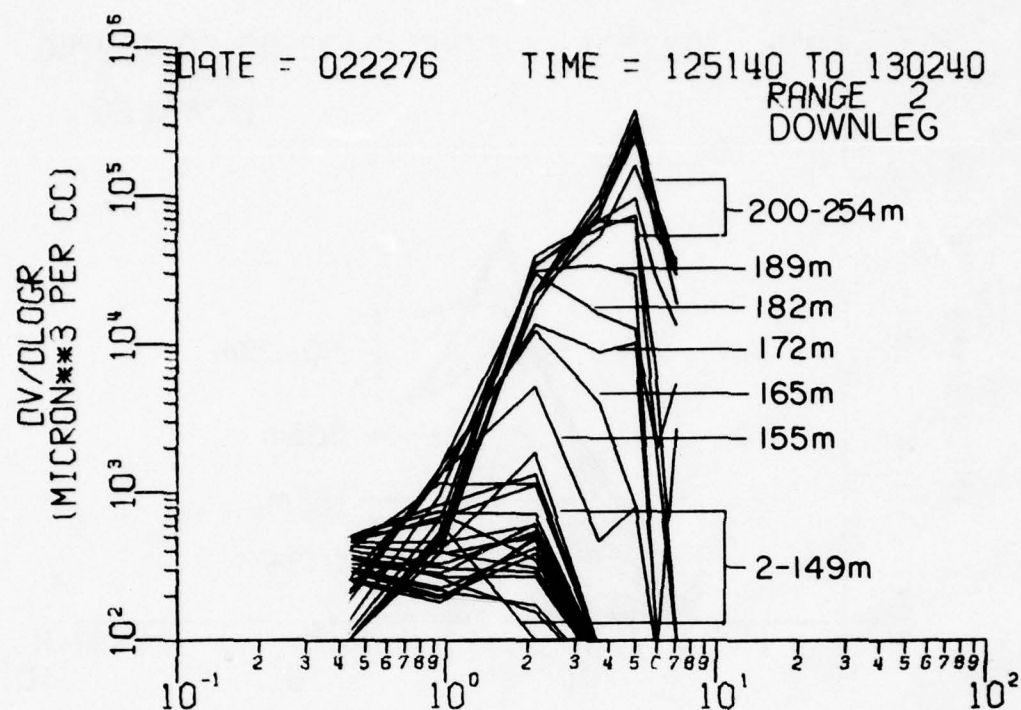


Figure A3-e

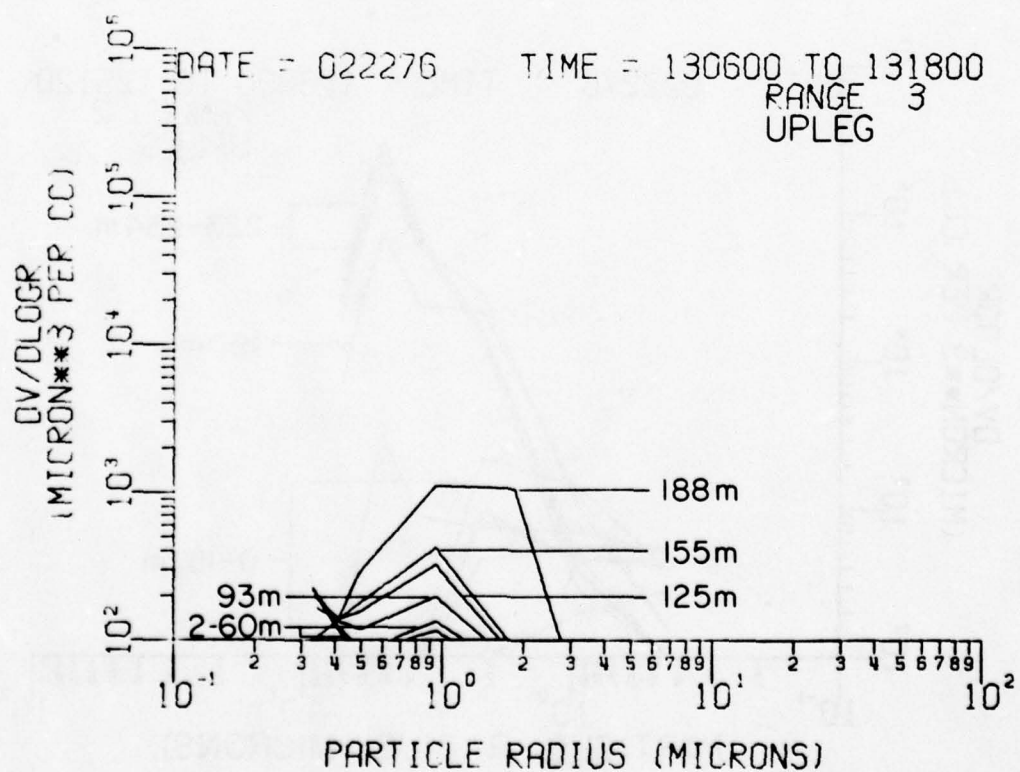
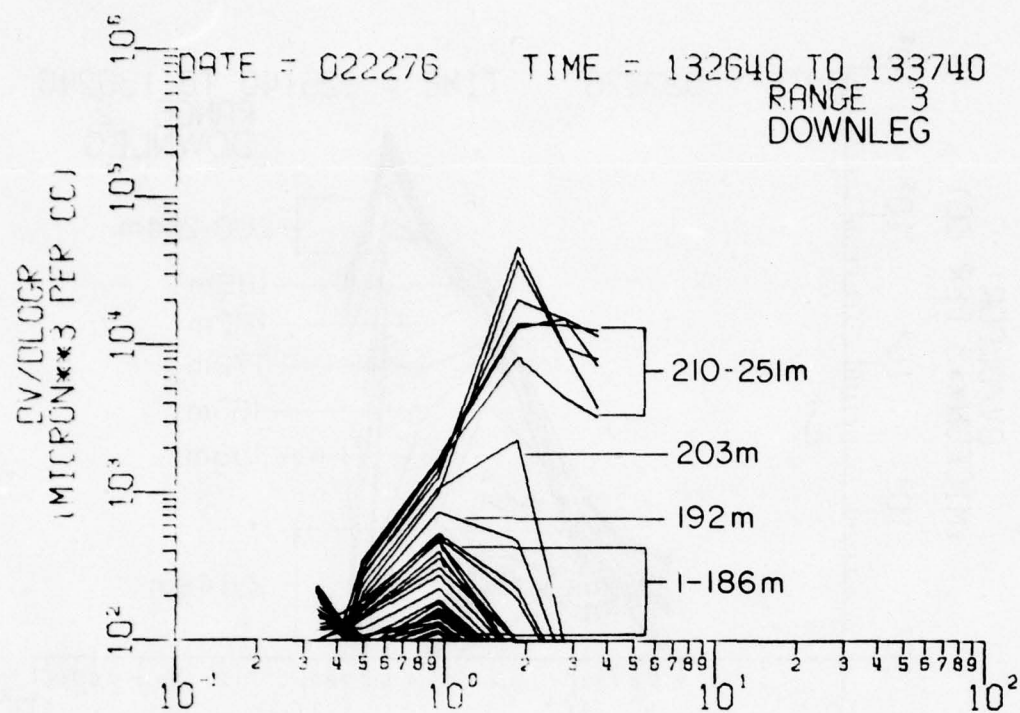


Figure A3-f

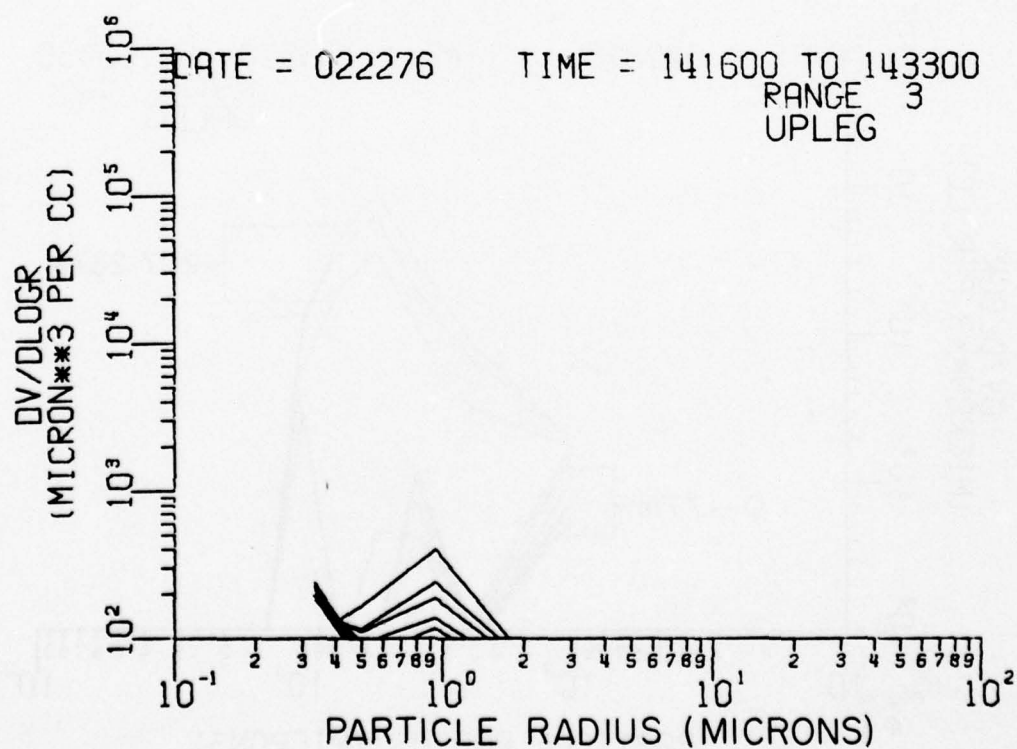
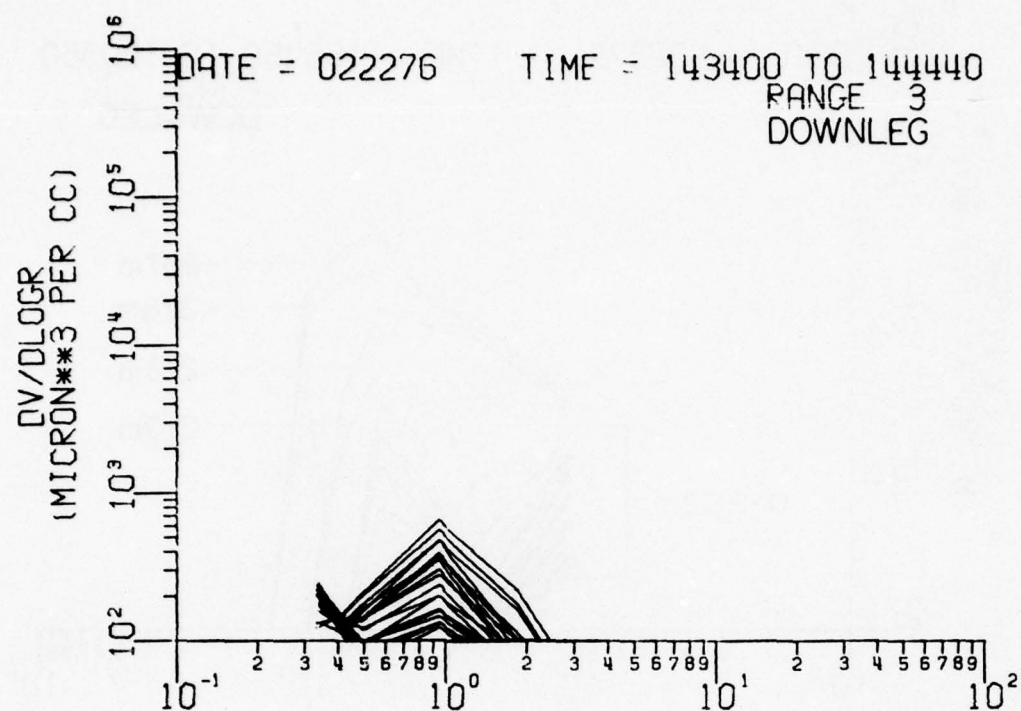


Figure A3-g

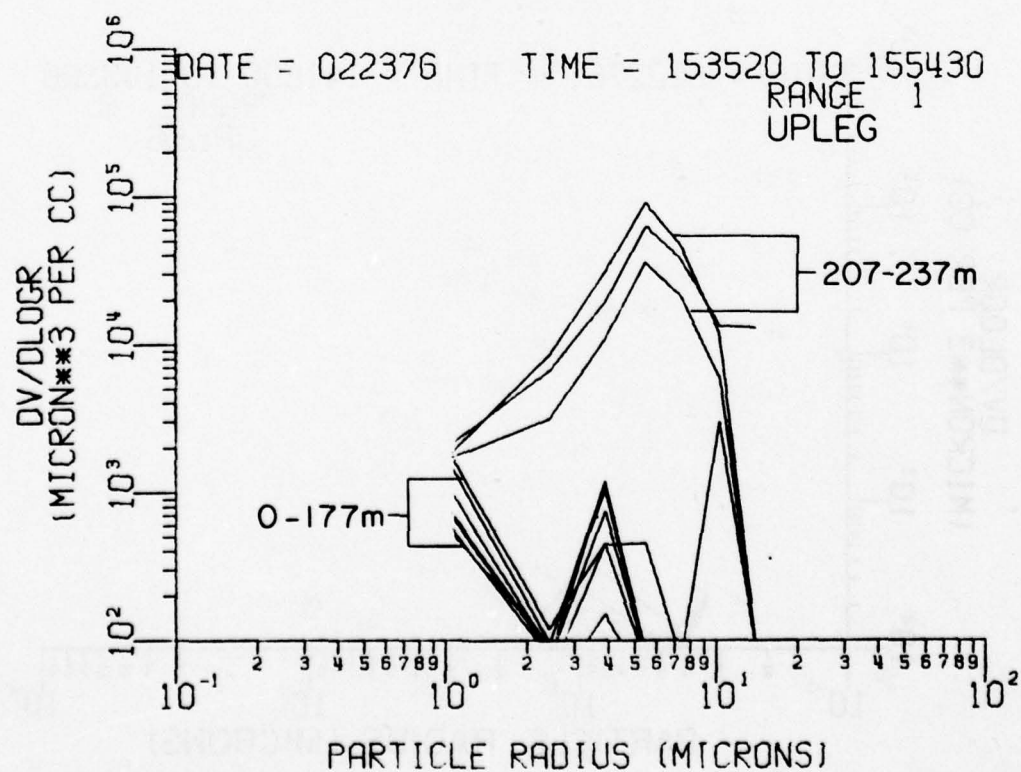
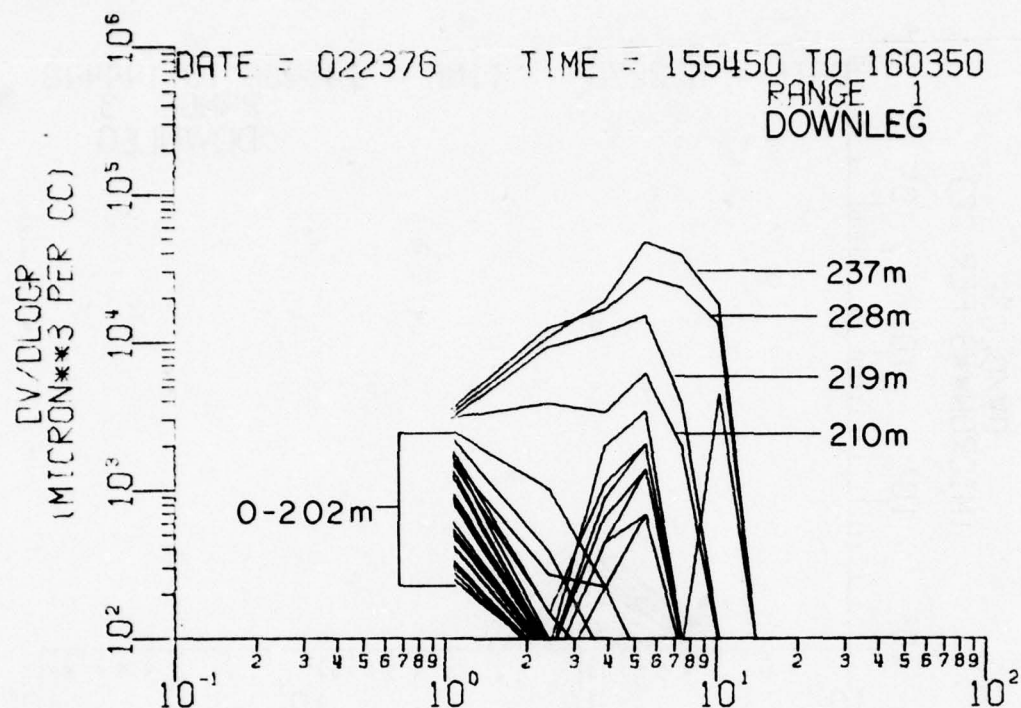


Figure A3-h

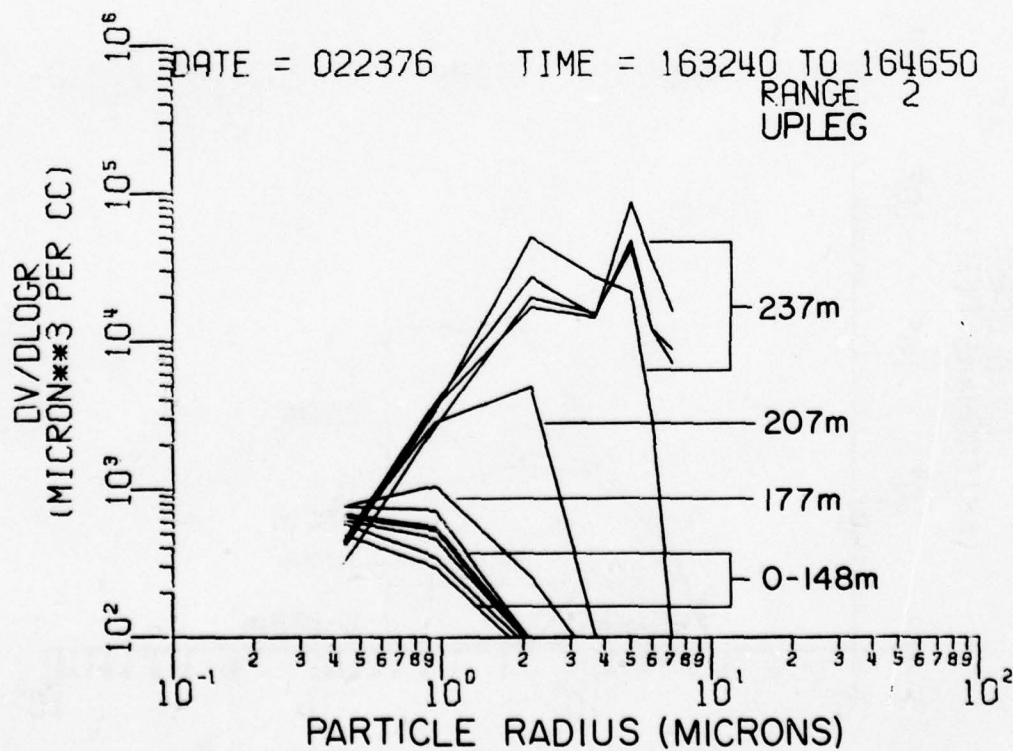
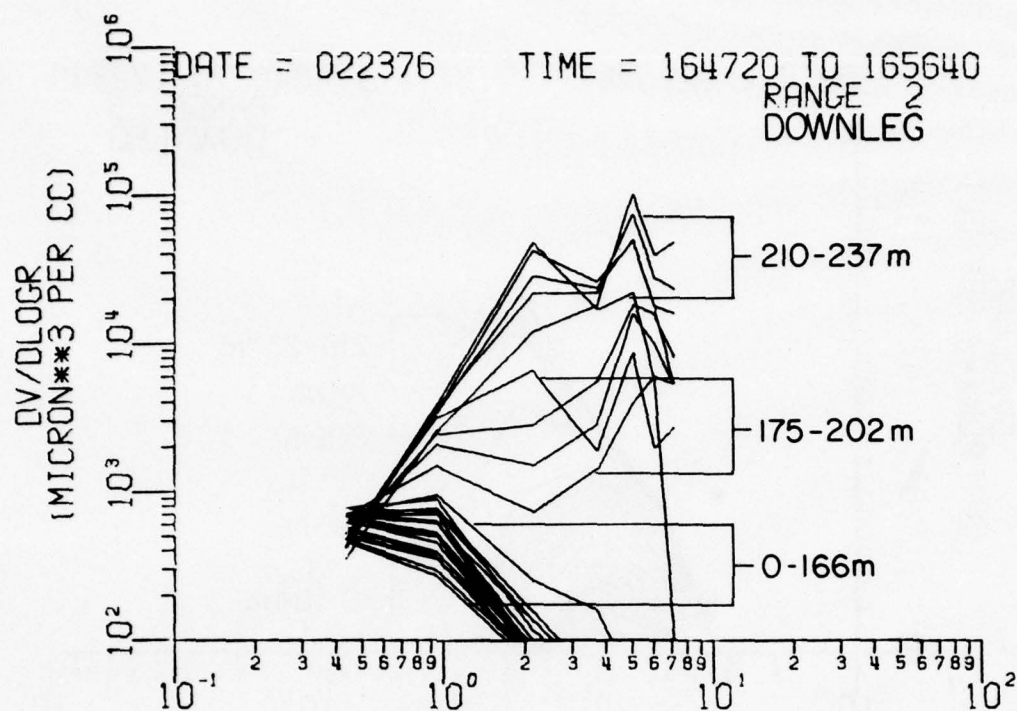


Figure A3-i

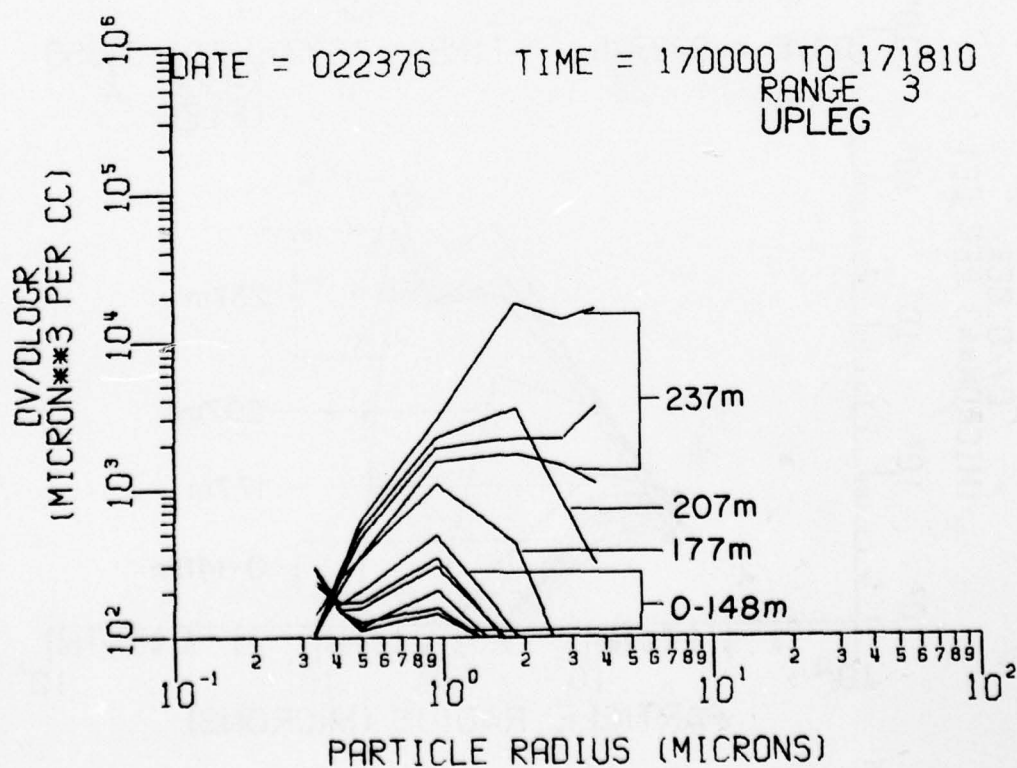
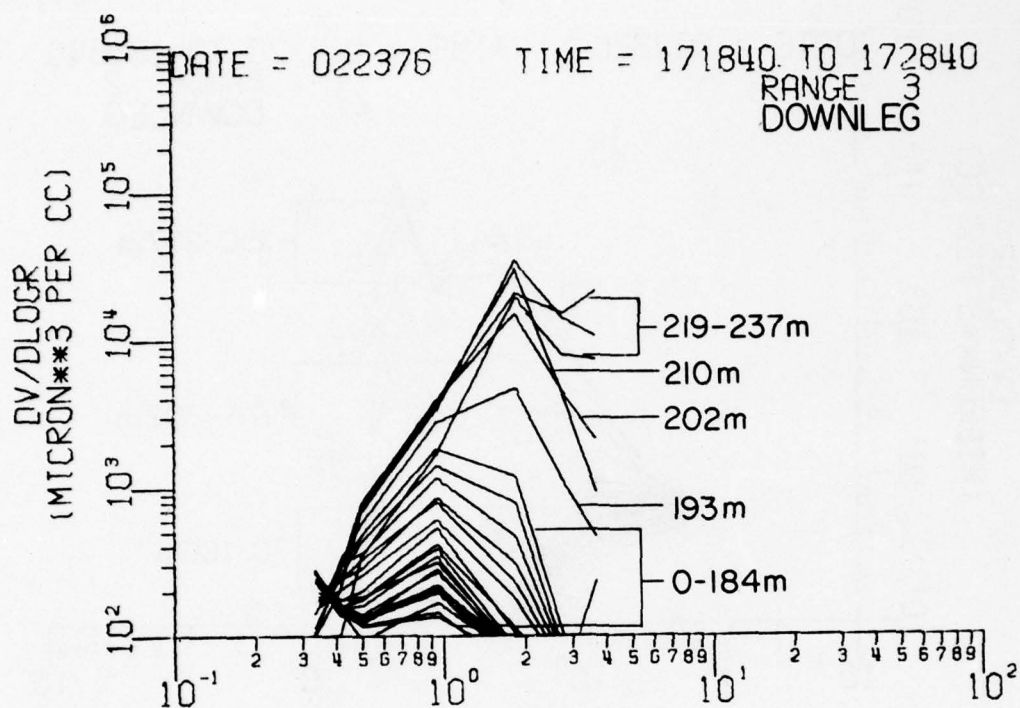


Figure A3-j

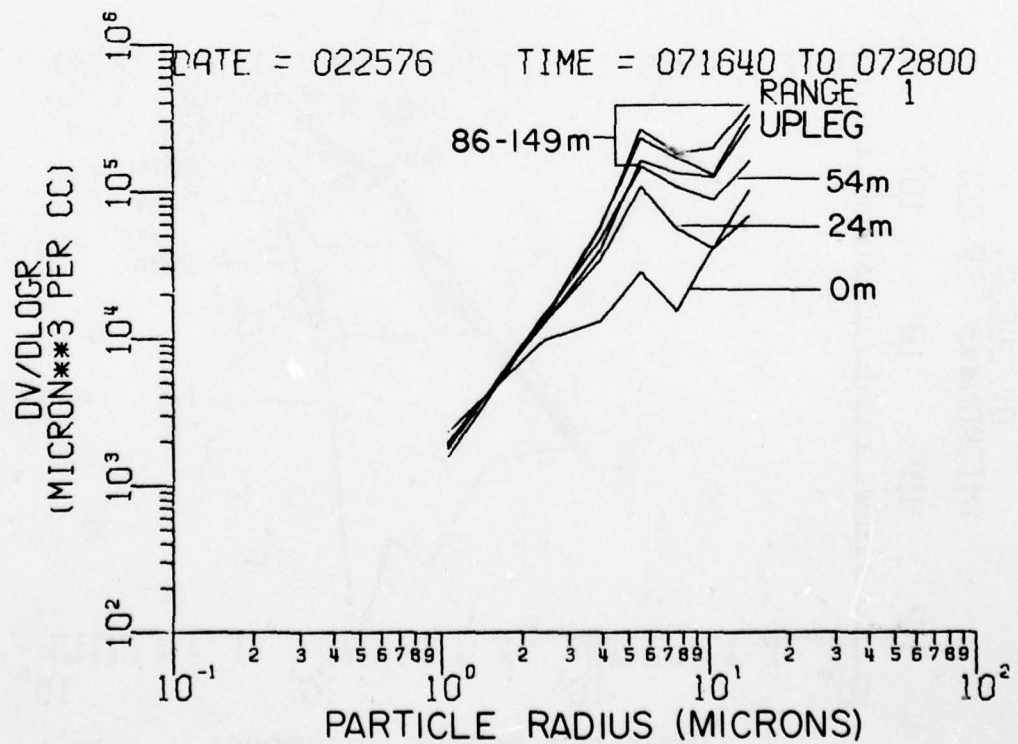
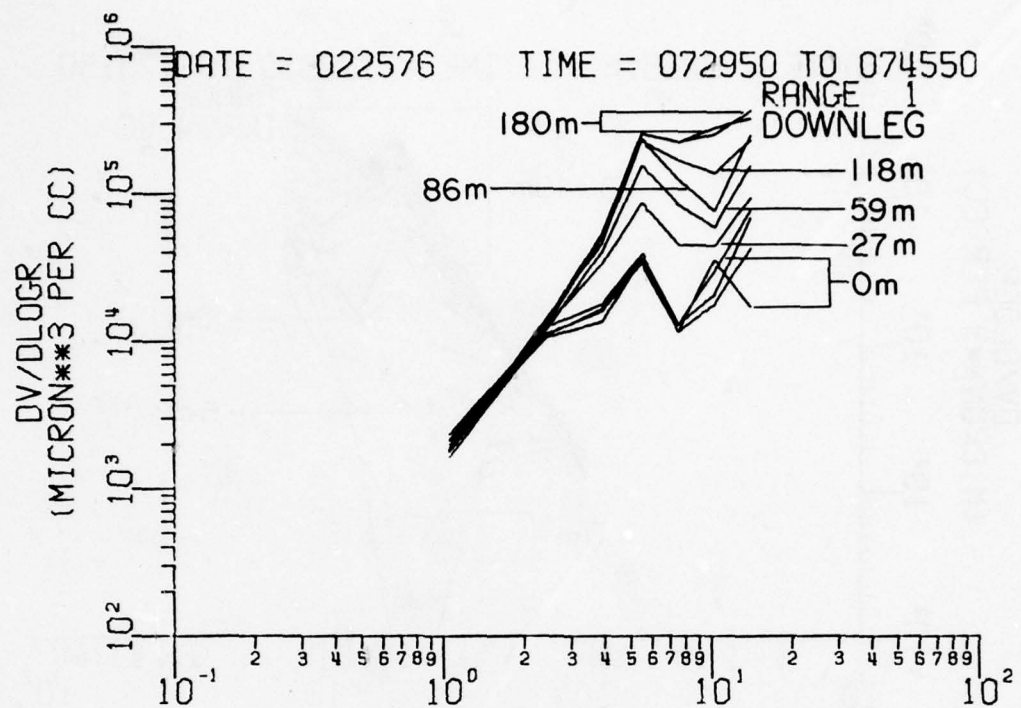


Figure A3-k

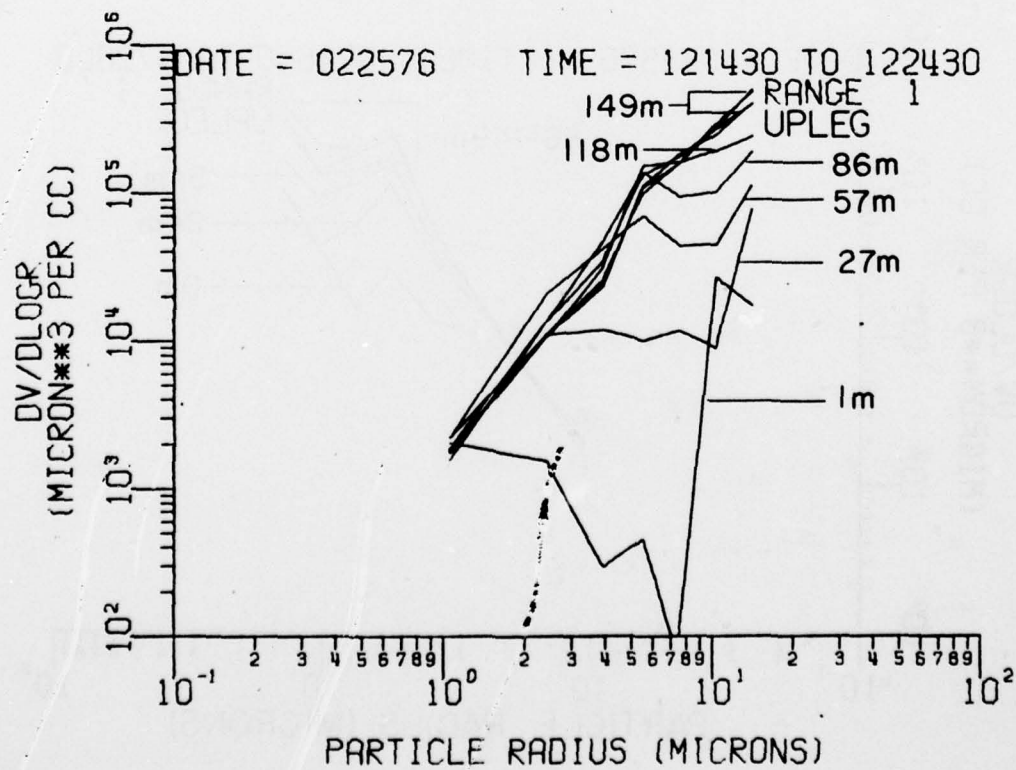
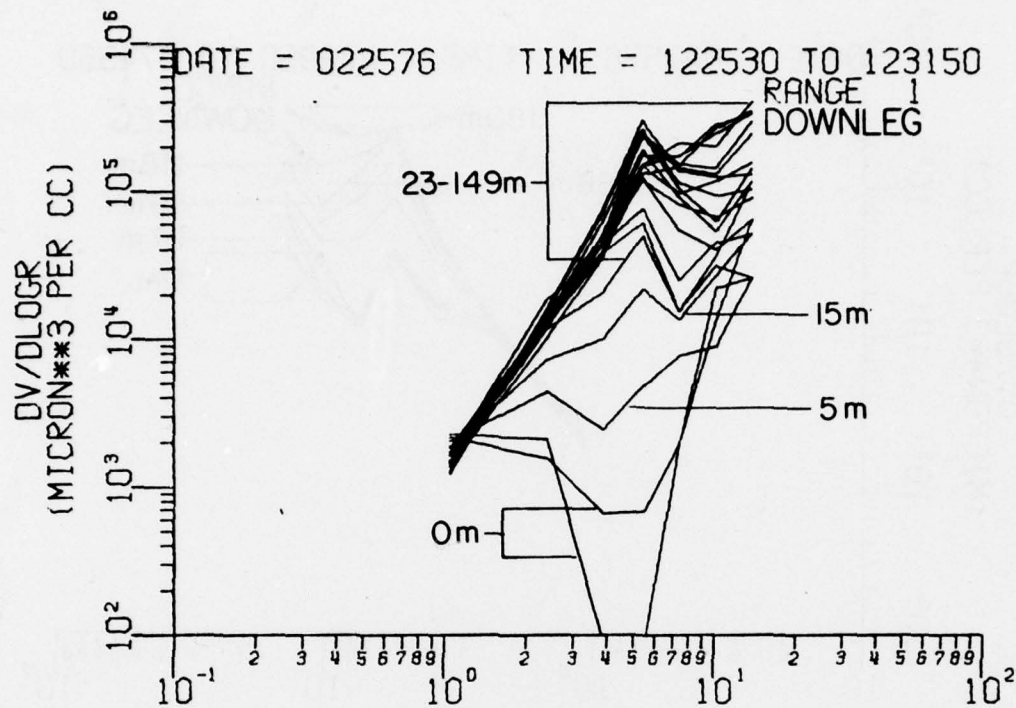


Figure A3-1

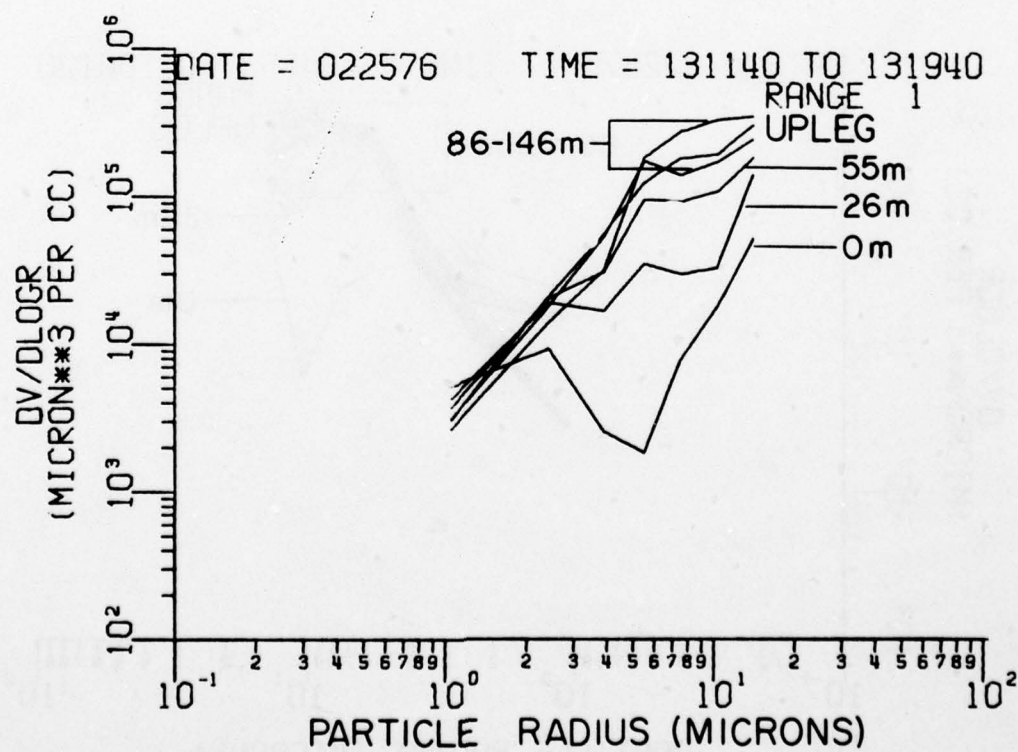
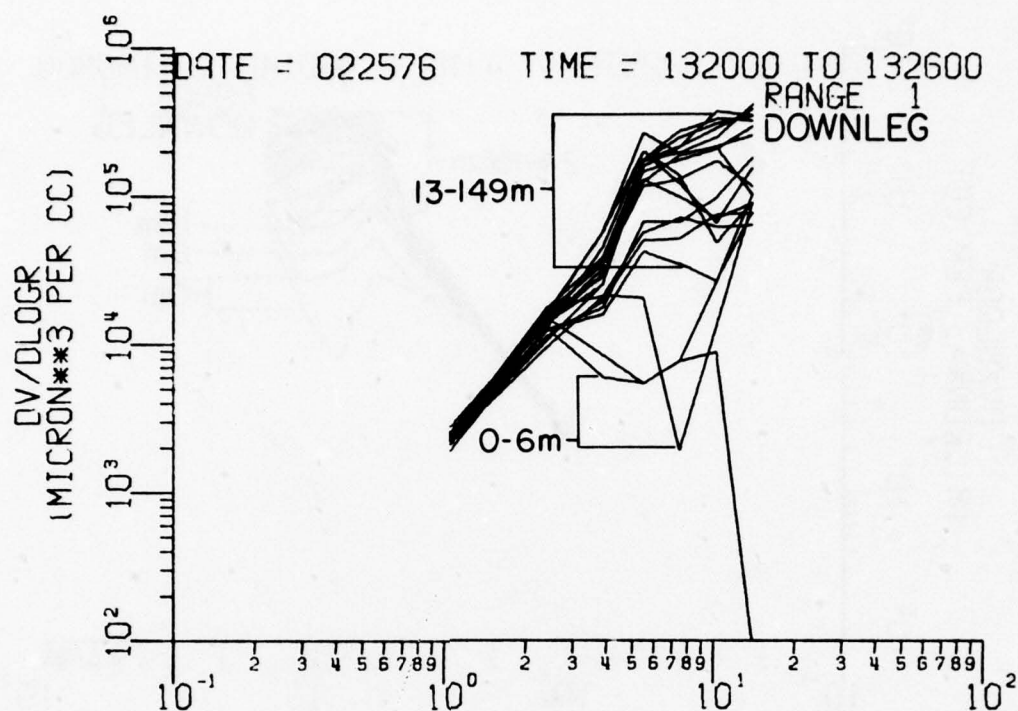


Figure A3-m

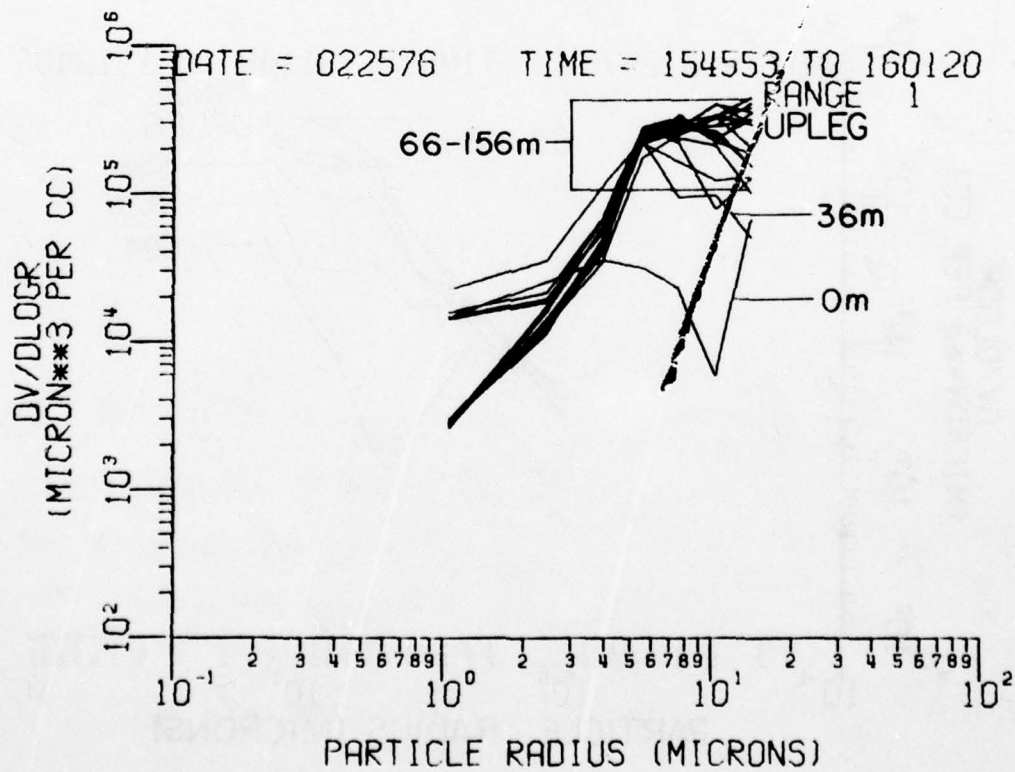
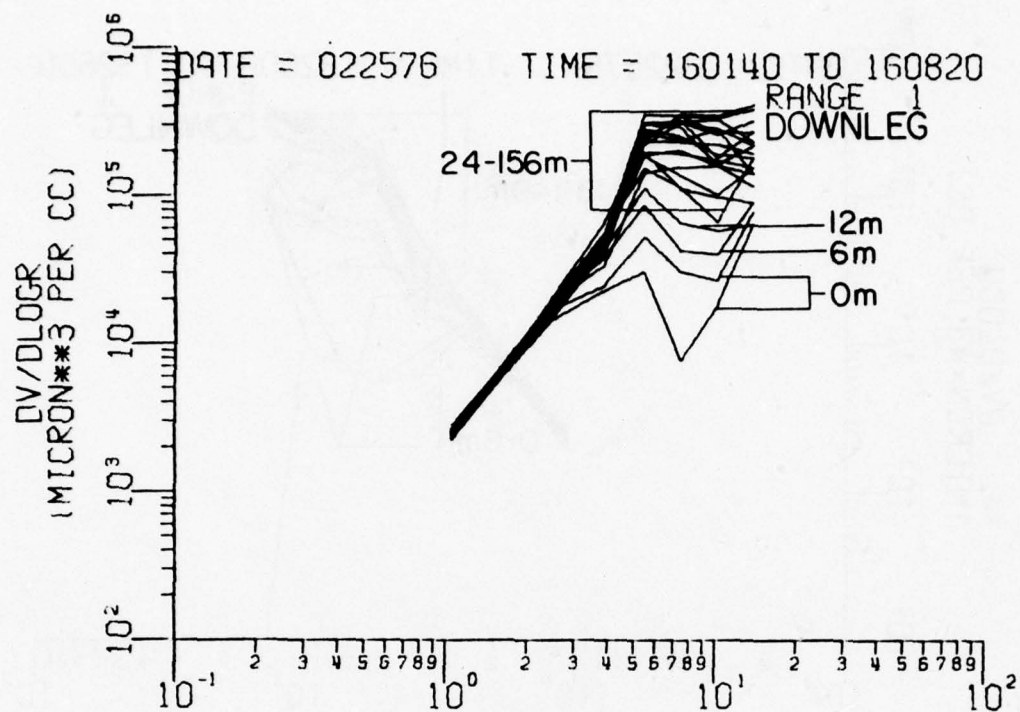


Figure A3-n

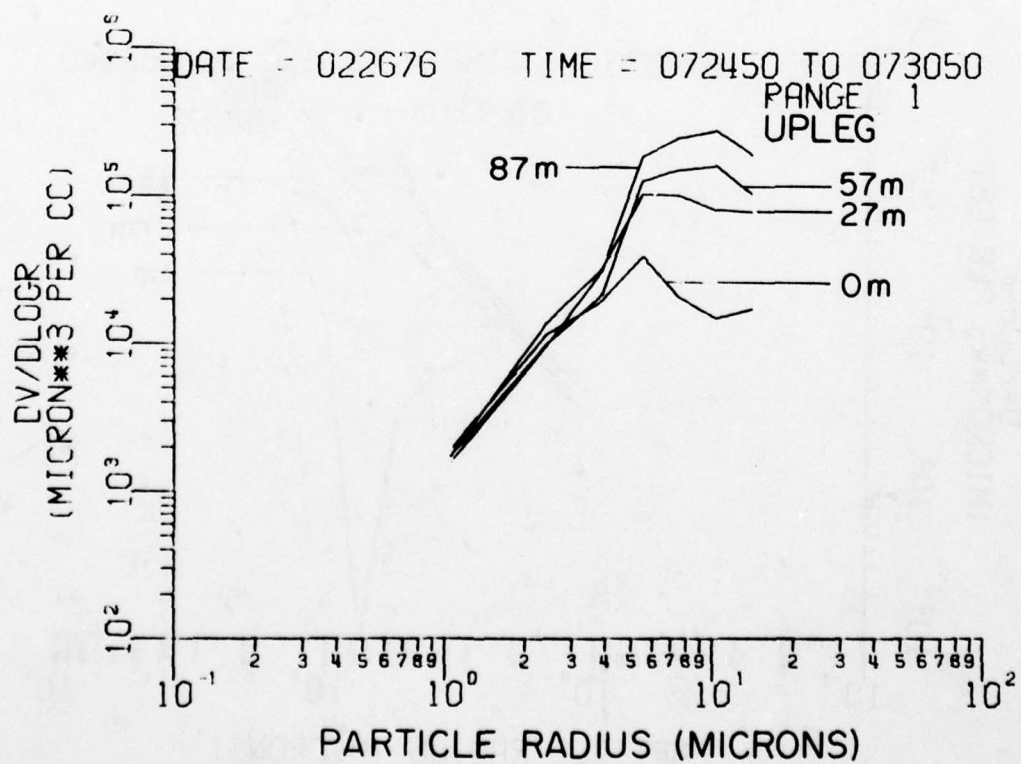
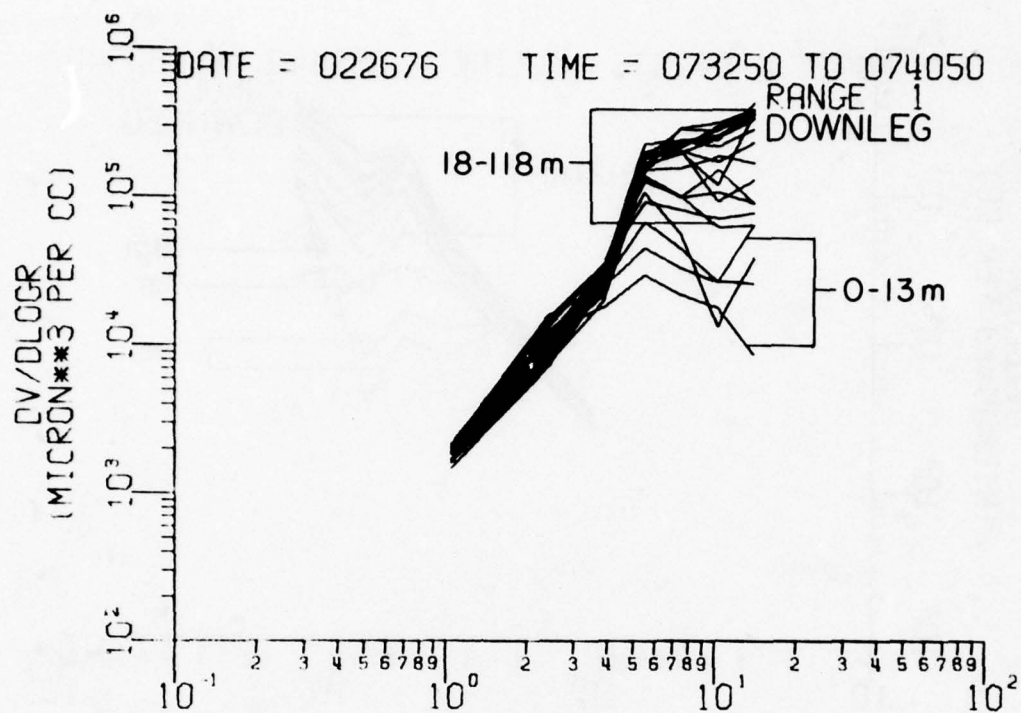


Figure A3-o

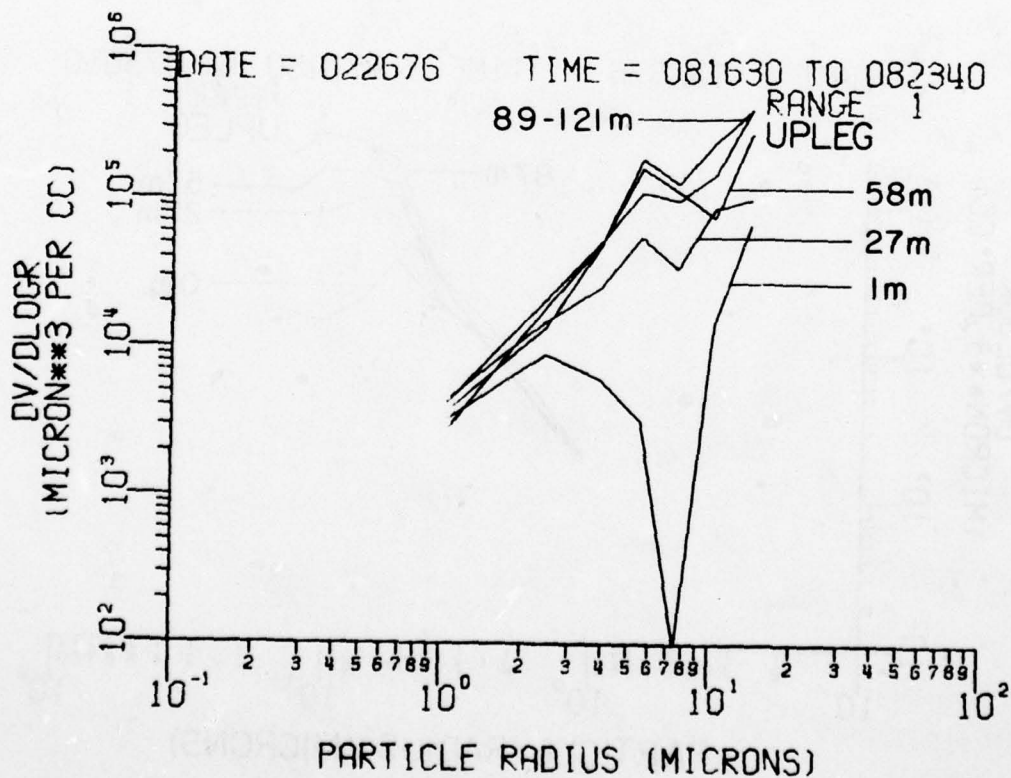
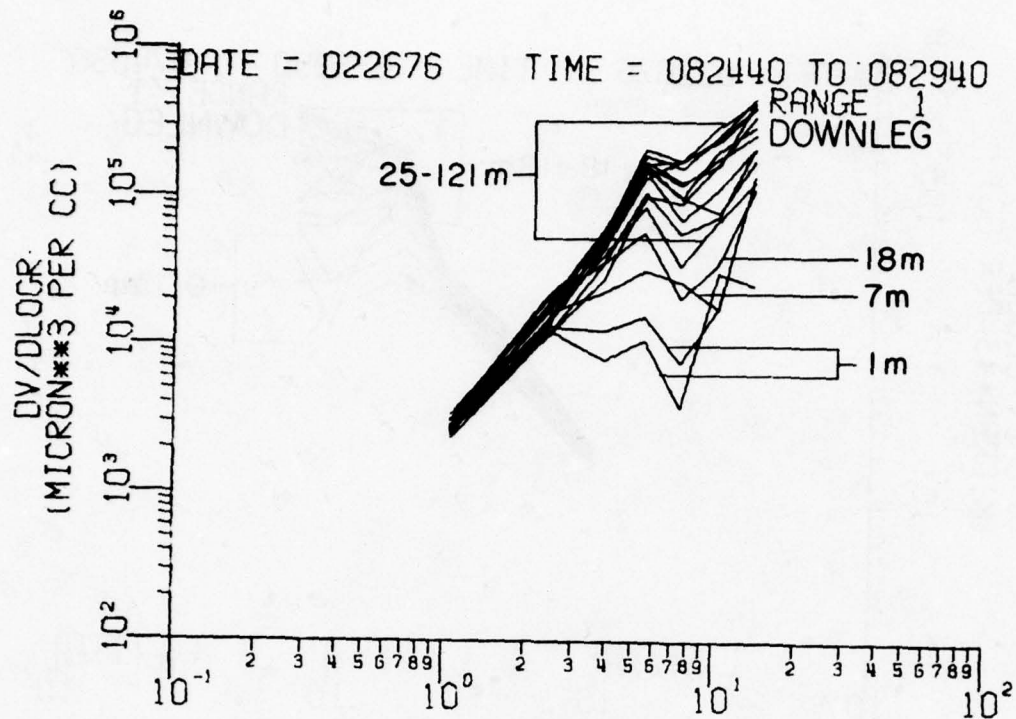


Figure A3-p

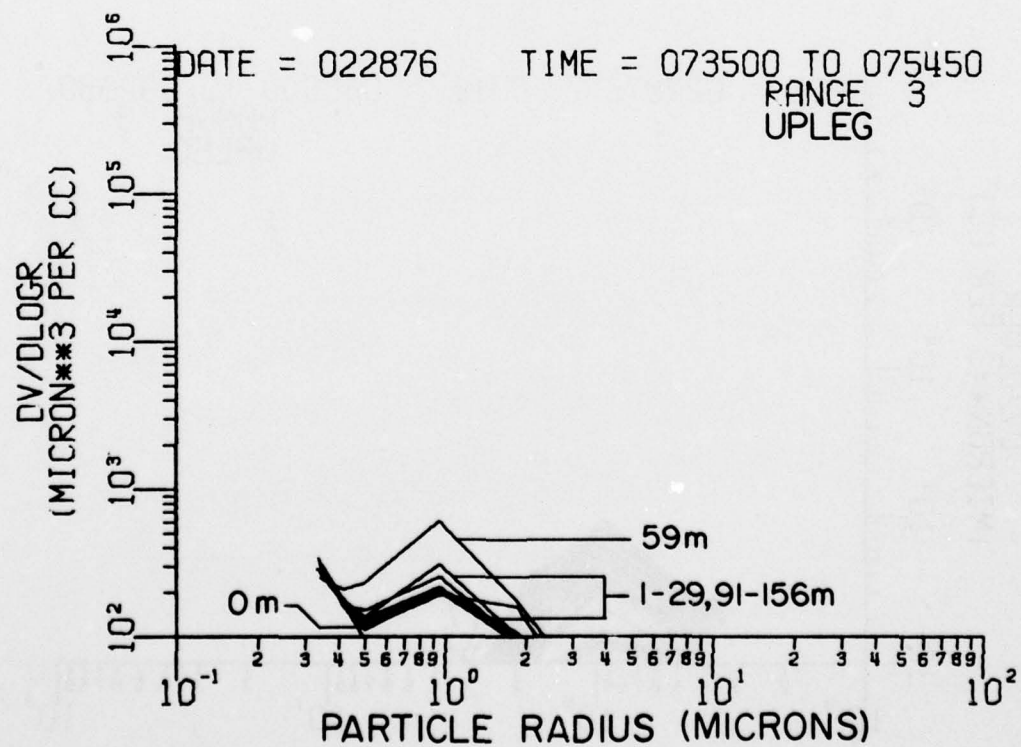
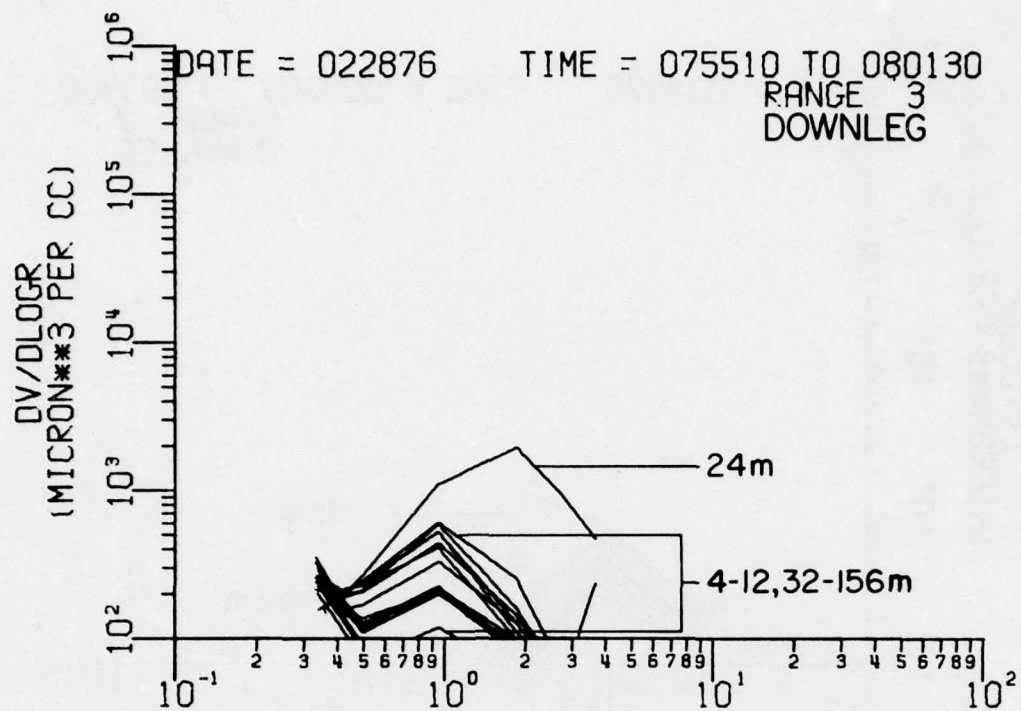


Figure A3-q

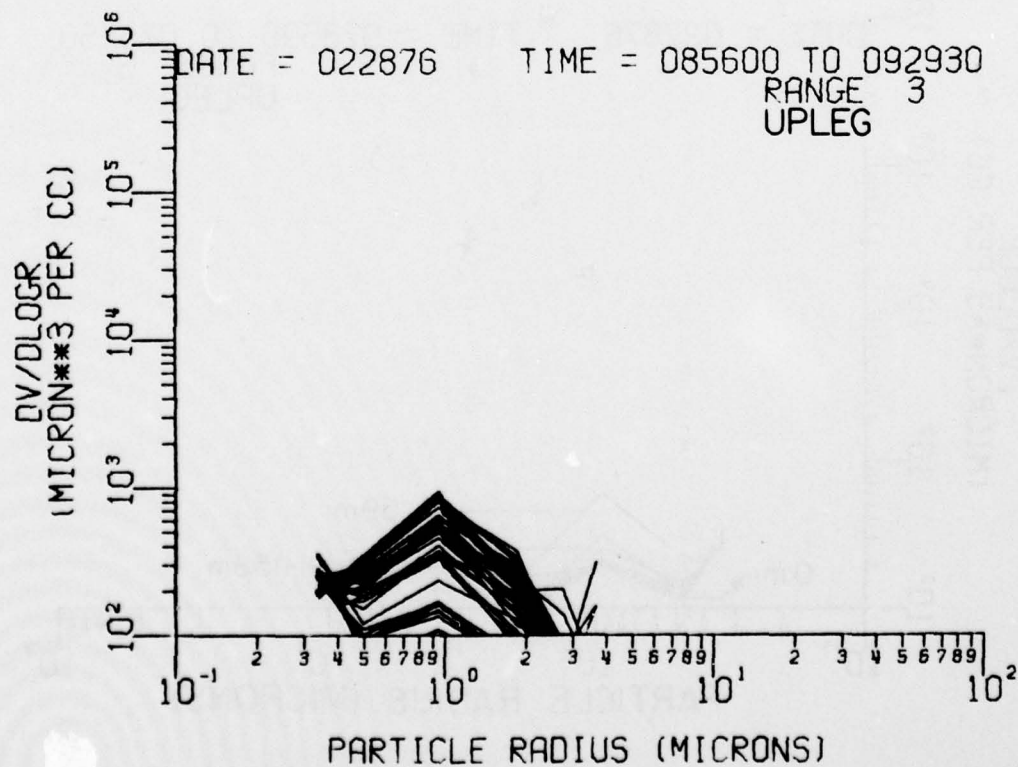
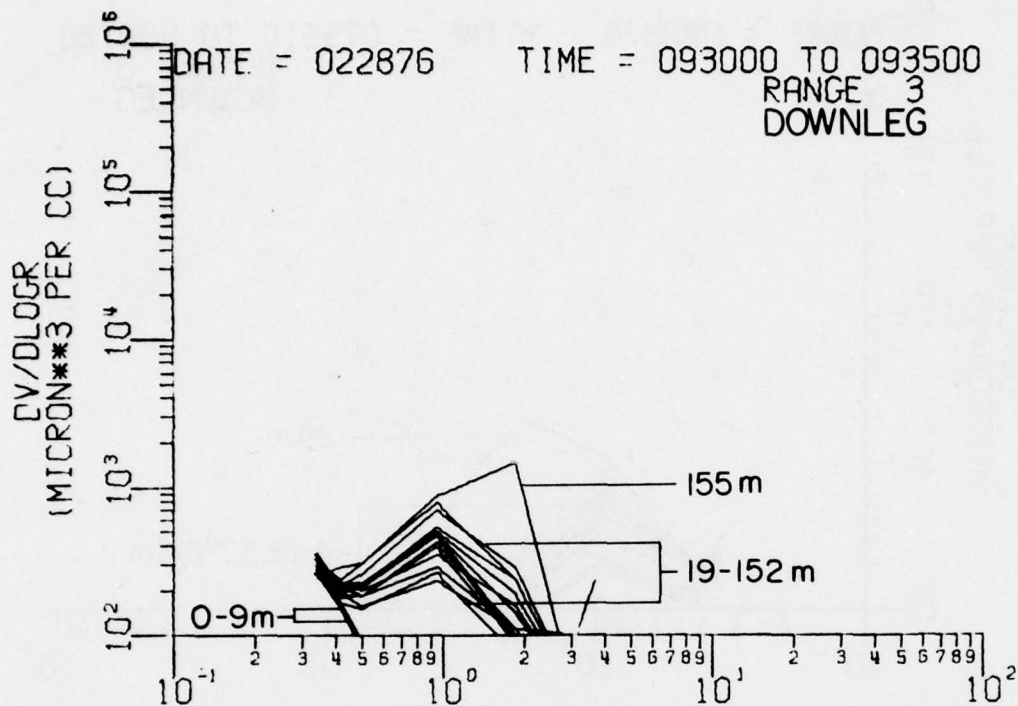


Figure A3-r

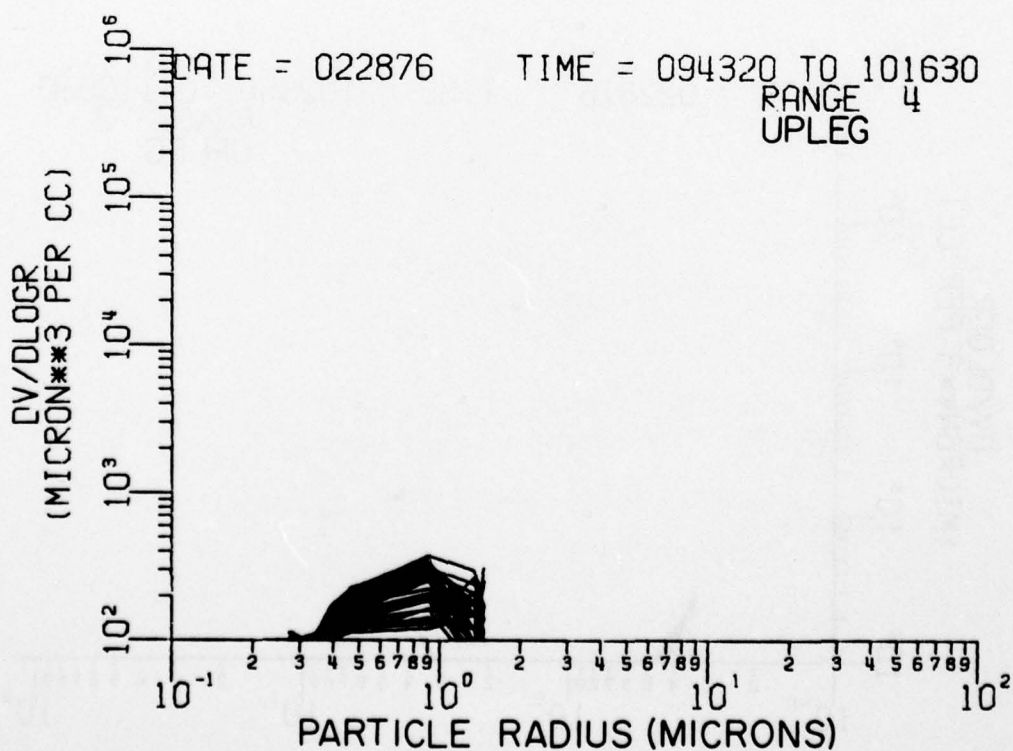
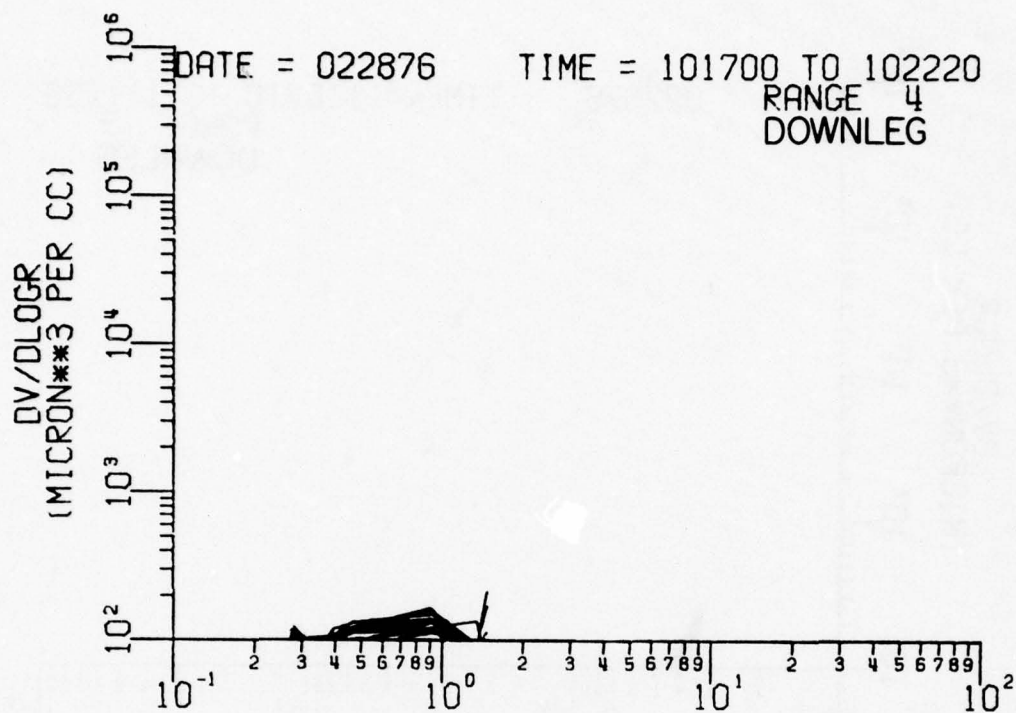


Figure A3-s

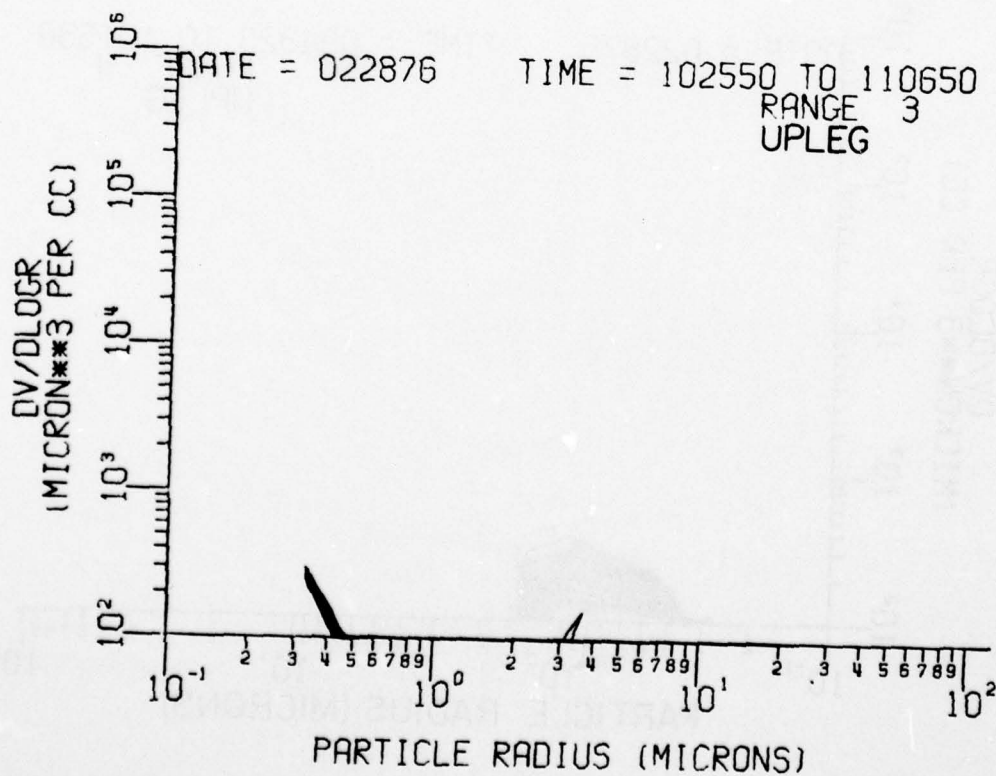
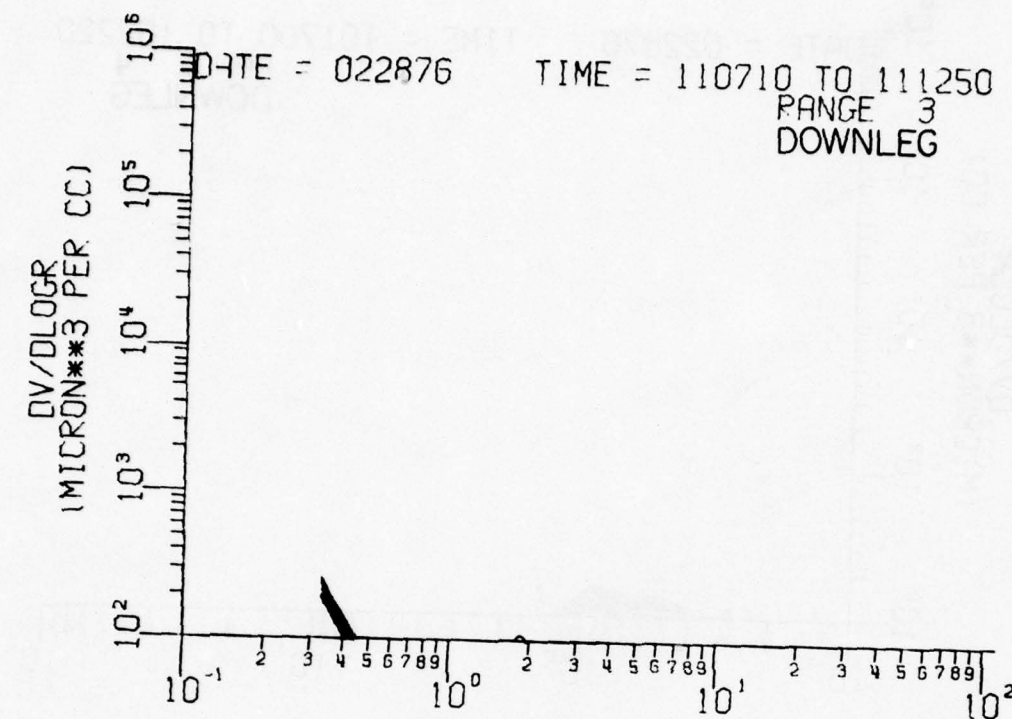


Figure A3-t

DATE = 12/14/75

TIME = 11:36:13 TO 12:33:33

LIQUID WATER CONTENT (GM PER METER**3)

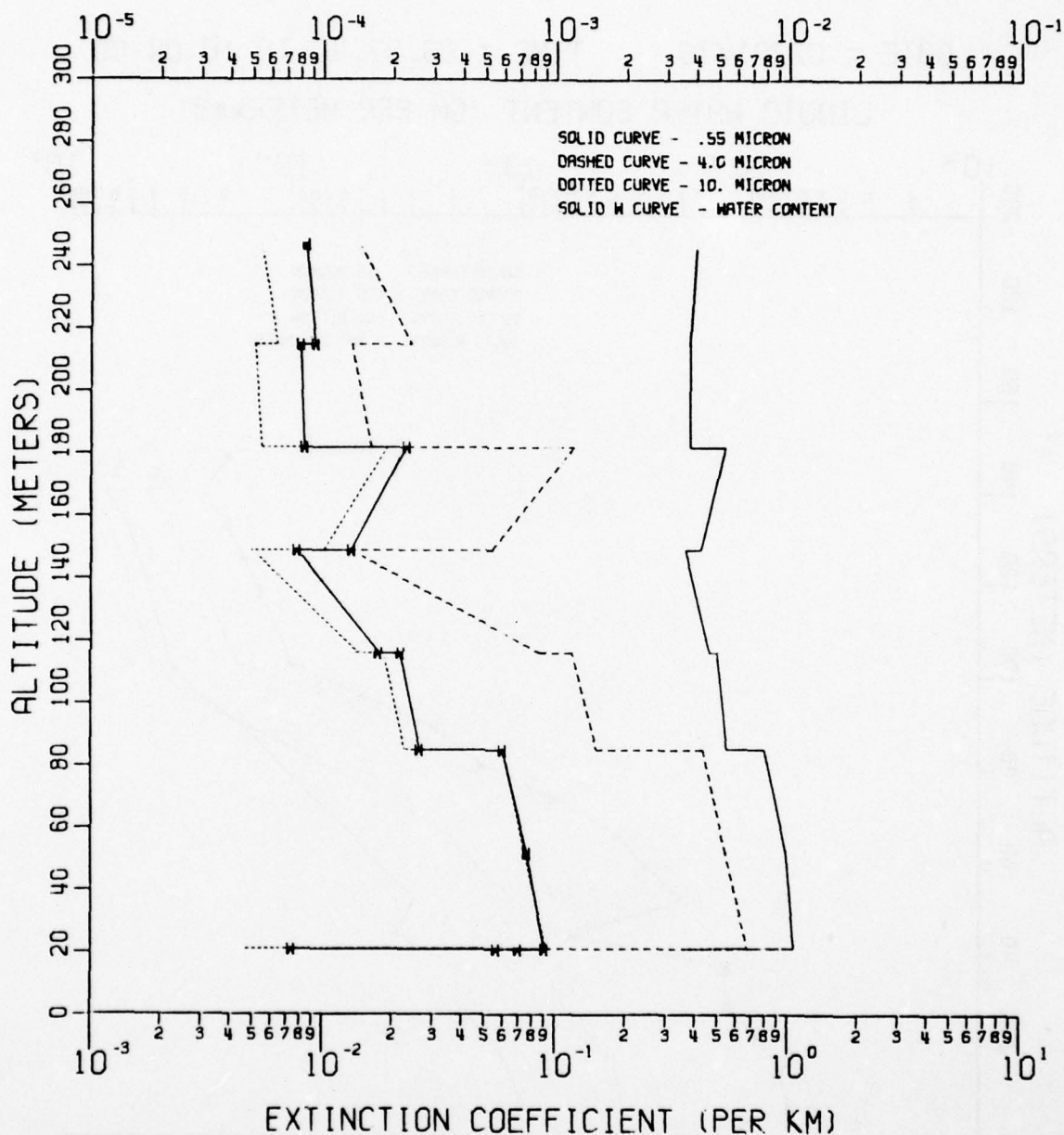


Figure A4-a. Profiles of particulate extinction and liquid water content inferred from particle size distribution measurements (shown in Fig. A1 made with a balloon-borne aerosol counter. Only the downleg portions of each traverse are presented. The solid "w" curve indicates liquid water content and the solid, dashed, and dotted curves indicate the inferred extinction at 0.55 μ m, 4 μ m, and 10 μ m wavelengths. The date, local time (hours, minutes, seconds) and time interval for the measurements are given.

DATE = 02/21/76

TIME = 09:57:40 TO 10:04:00

LIQUID WATER CONTENT (GM PER METER**3)

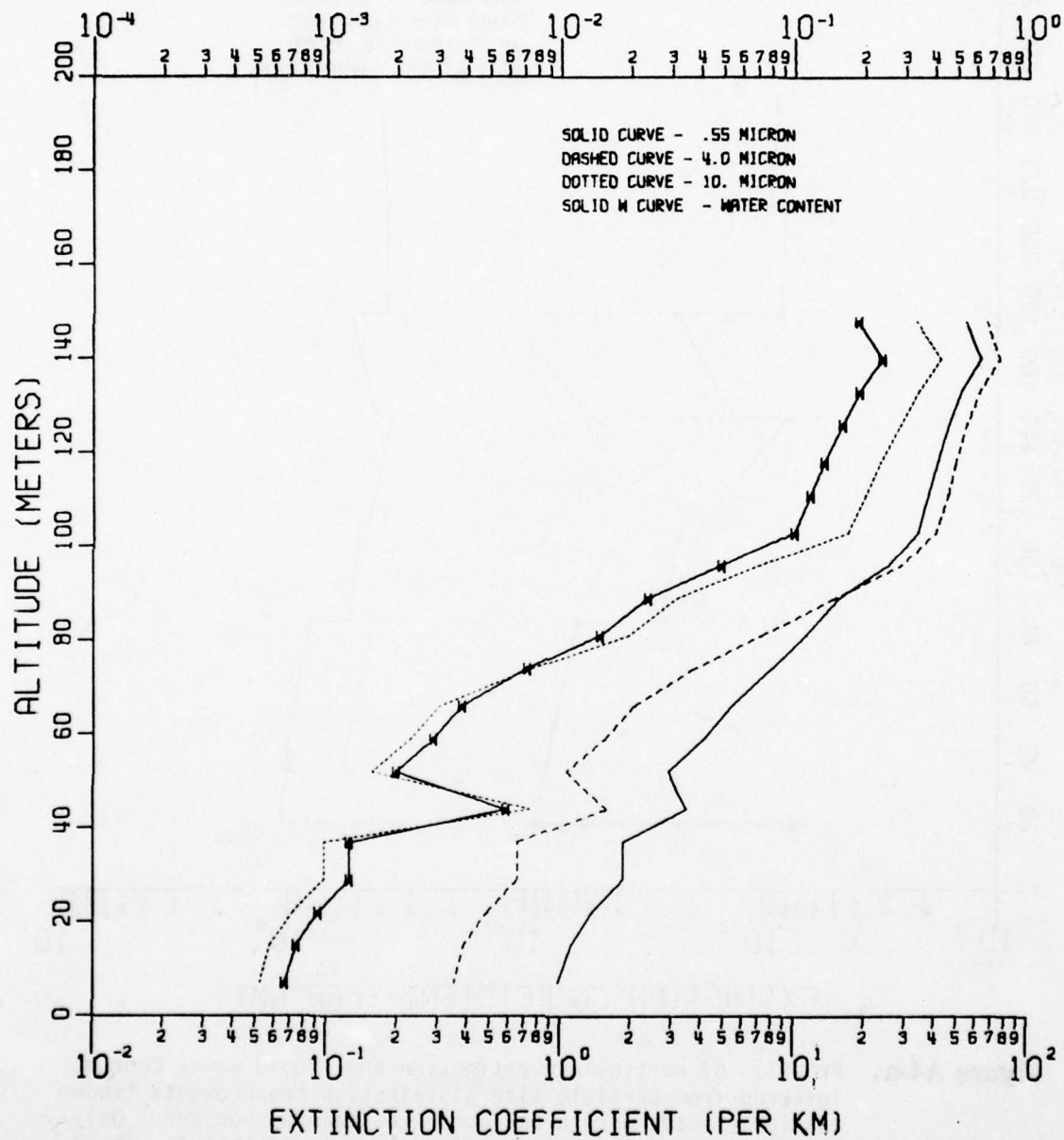


Figure A4-b

DATE = 02/21/76

TIME = 10:39:20 TO 10:46:10

LIQUID WATER CONTENT (GM PER METER**3)

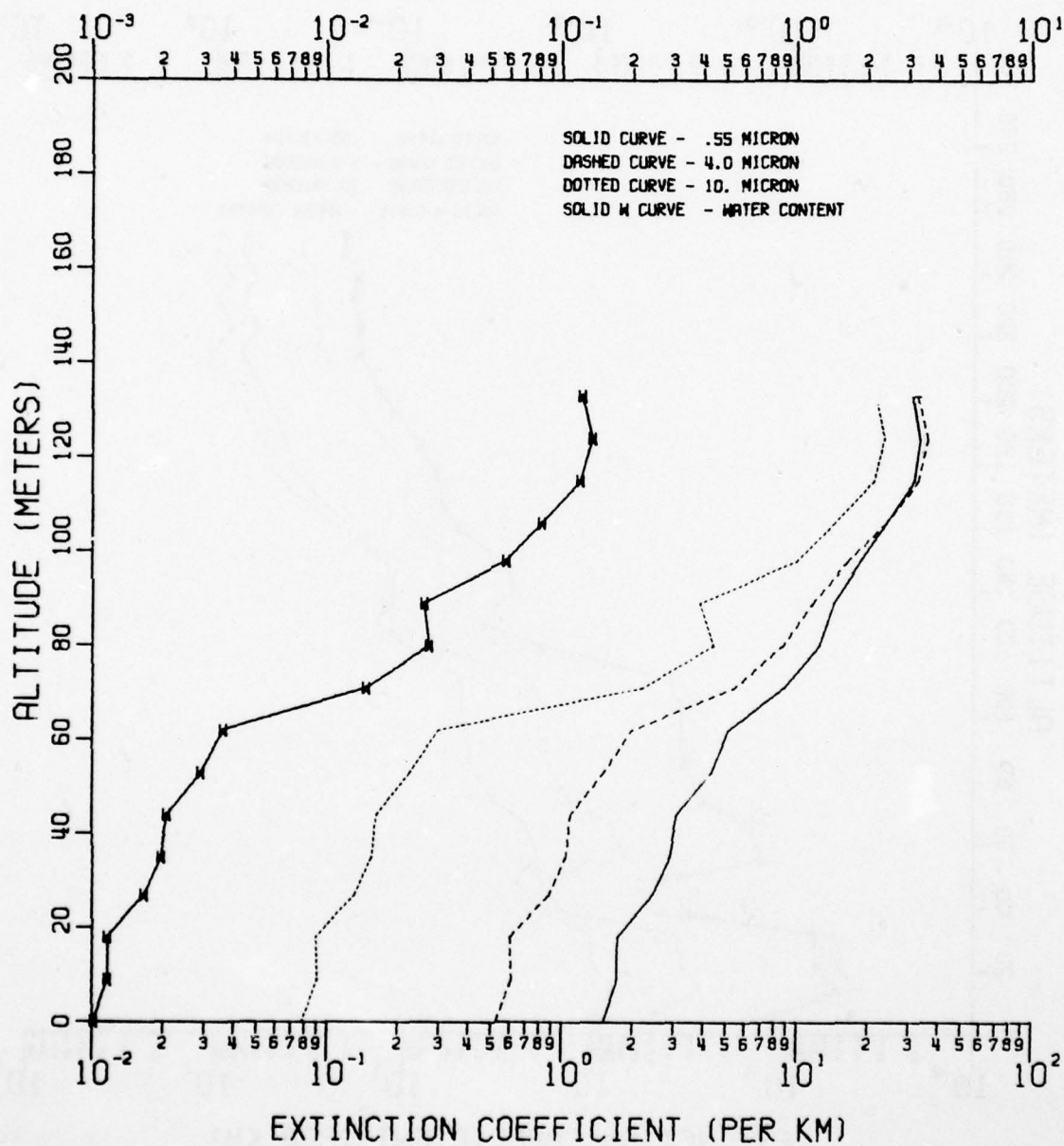


Figure A4-c

DATE = 02/22/76

TIME = 12:16:50 TO 12:26:30

LIQUID WATER CONTENT (GM PER METER**3)

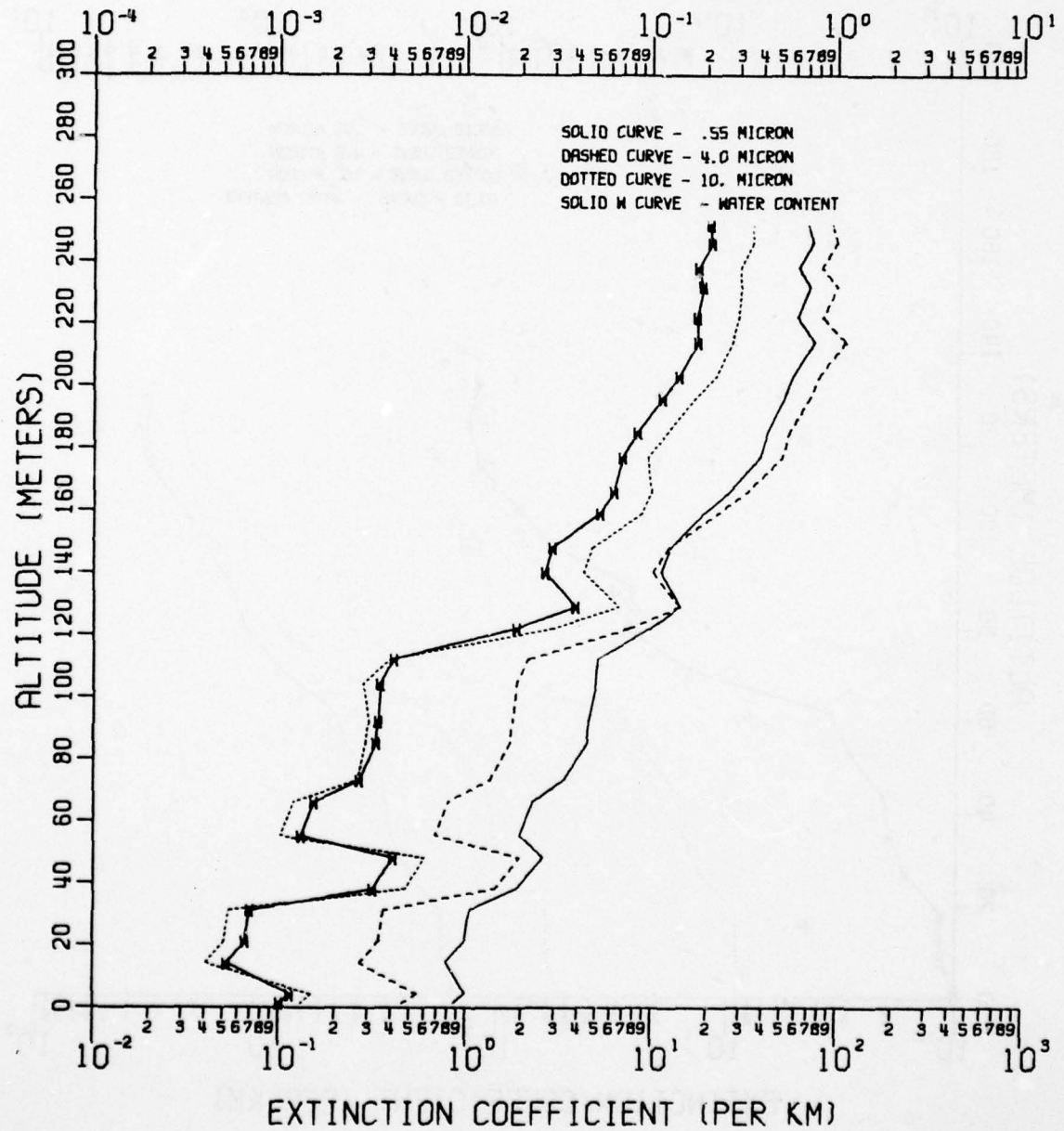


Figure A4-d

A-64

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DATE = 02/22/76

TIME = 12:51:40 TO 13:02:40

LIQUID WATER CONTENT (GM PER METER**3)

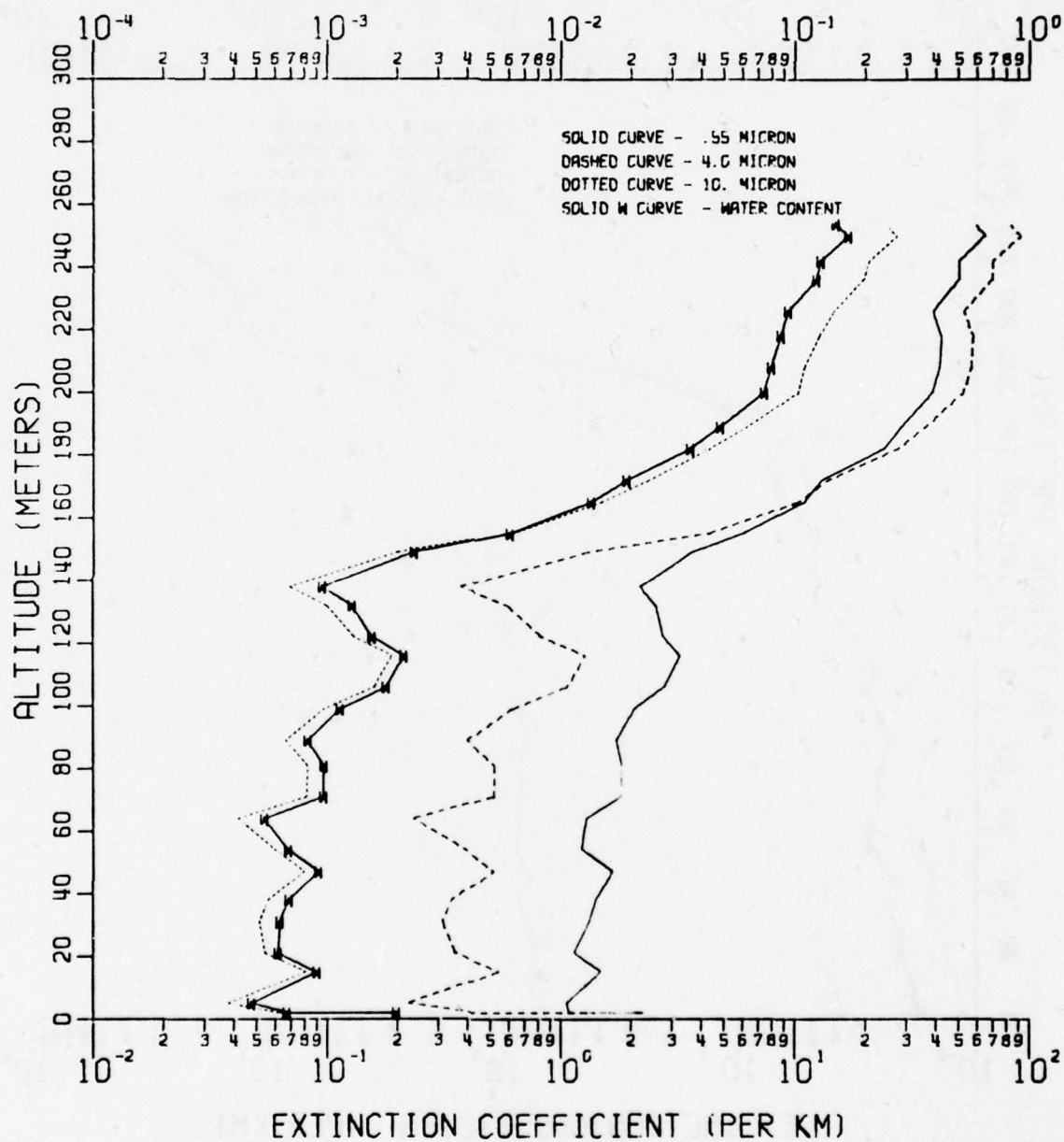


Figure A4-e

DATE = 02/22/76

TIME = 13:26:40 TO 13:37:40

LIQUID WATER CONTENT (GM PER METER**3)

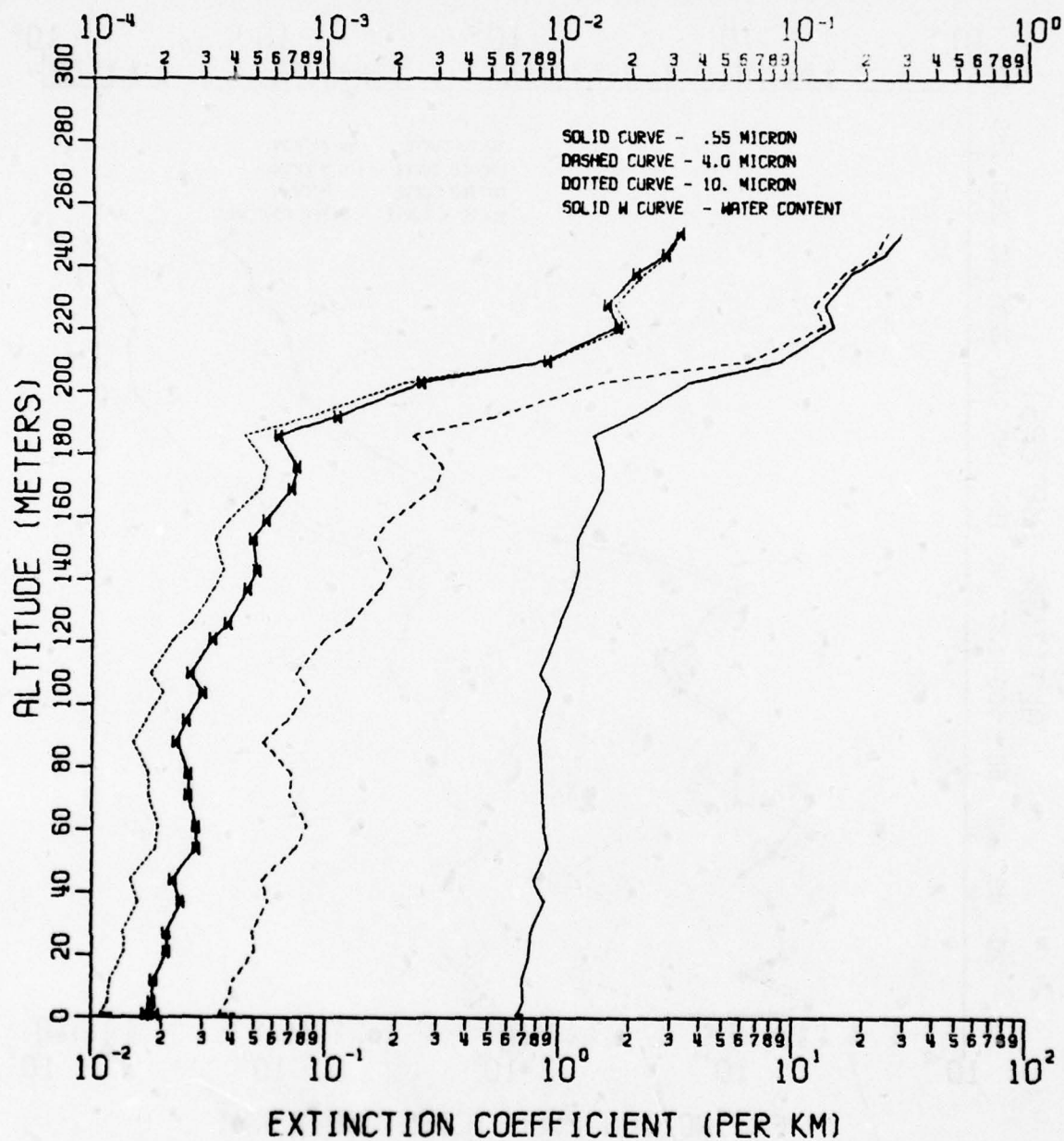


Figure A4-f

DATE = 02/22/76

TIME = 14:34:00 TO 14:44:40

LIQUID WATER CONTENT (GM PER METER**3)

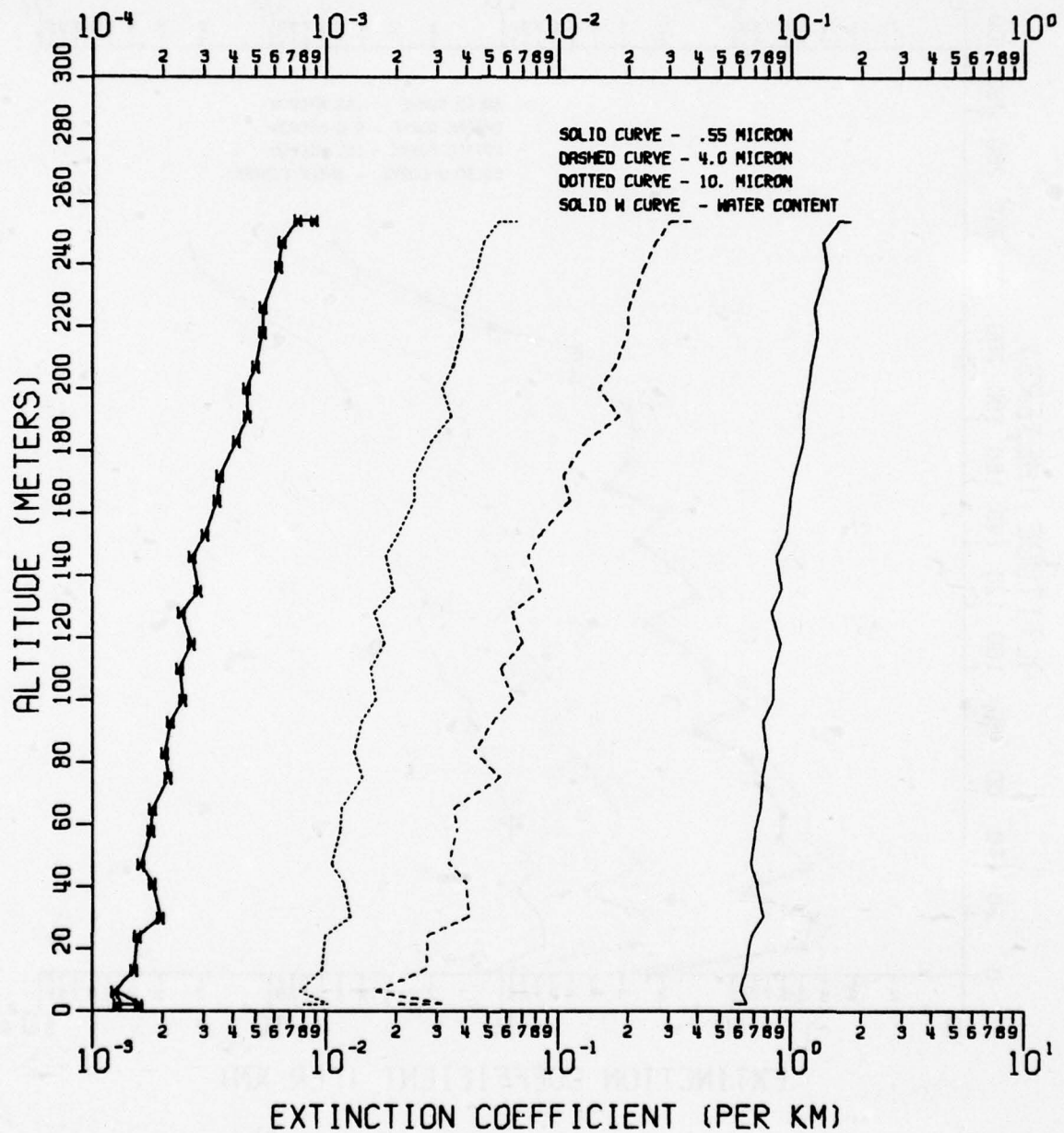


Figure A4-g

DATE = 02/23/76

TIME = 15:54:50 TO 16:03:50

LIQUID WATER CONTENT (GM PER METER**3)

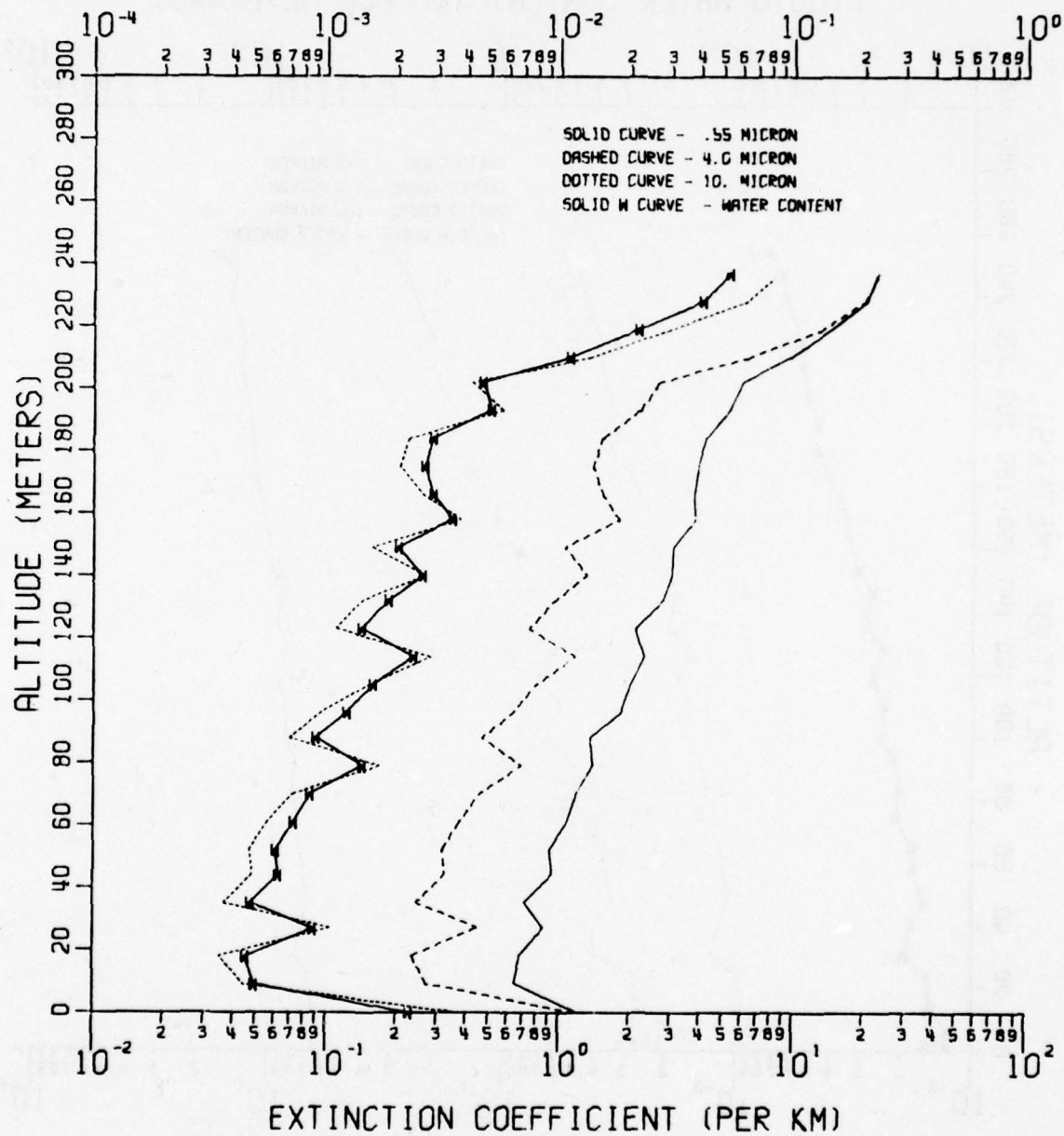


Figure A4-h

A-68

DATE = 02/23/76

TIME = 16:47:20 TO 16:56:40

LIQUID WATER CONTENT (GM PER METER**3)

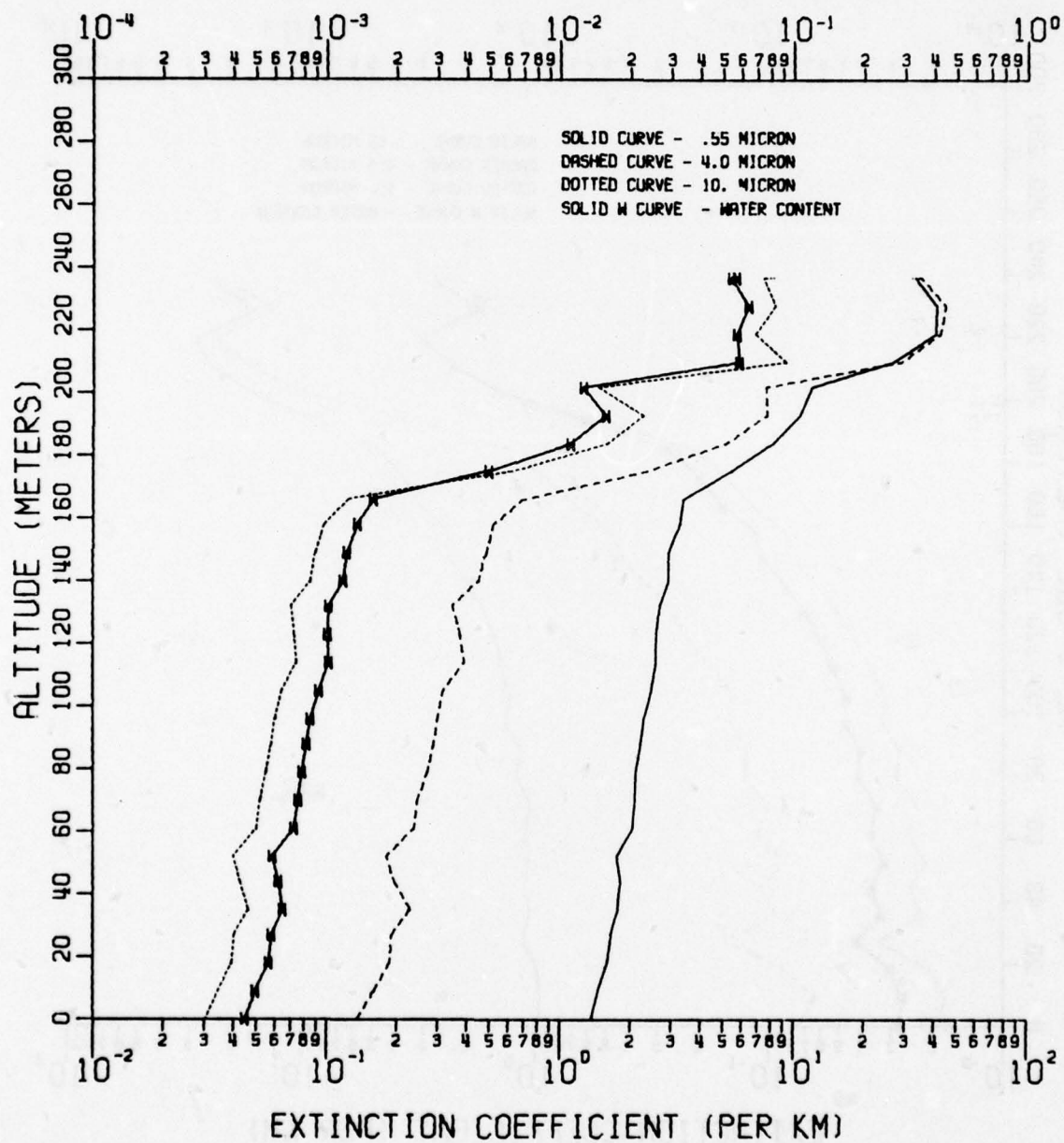


Figure A4-i

DATE = 02/23/76

TIME = 17:18:40 TO 17:28:40

LIQUID WATER CONTENT (GM PER METER**3)

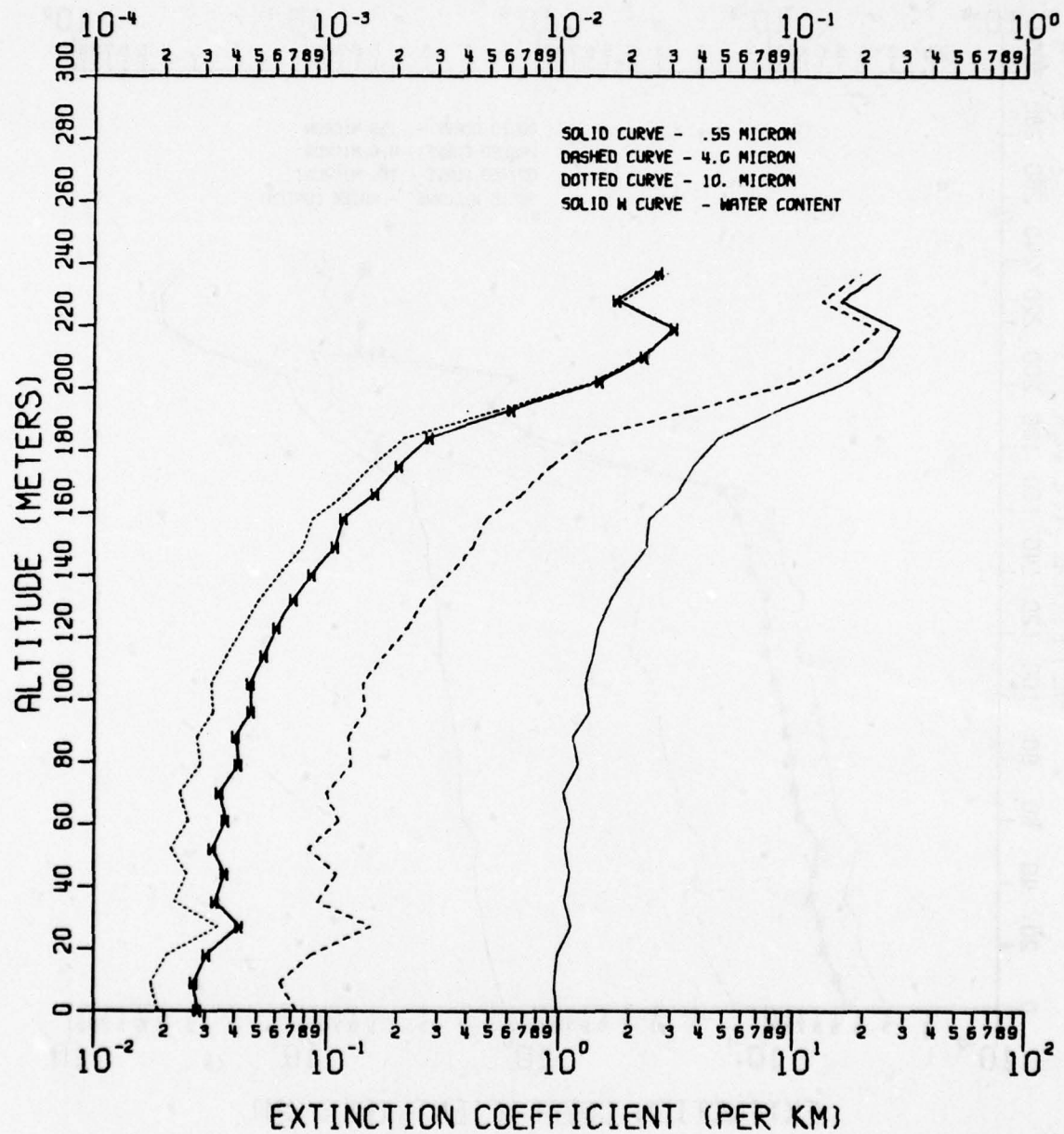


Figure A4-j

A-70

DATE = 02/25/76

TIME = 07:29:50 TO 07:45:50

LIQUID WATER CONTENT (GM PER METER**3)

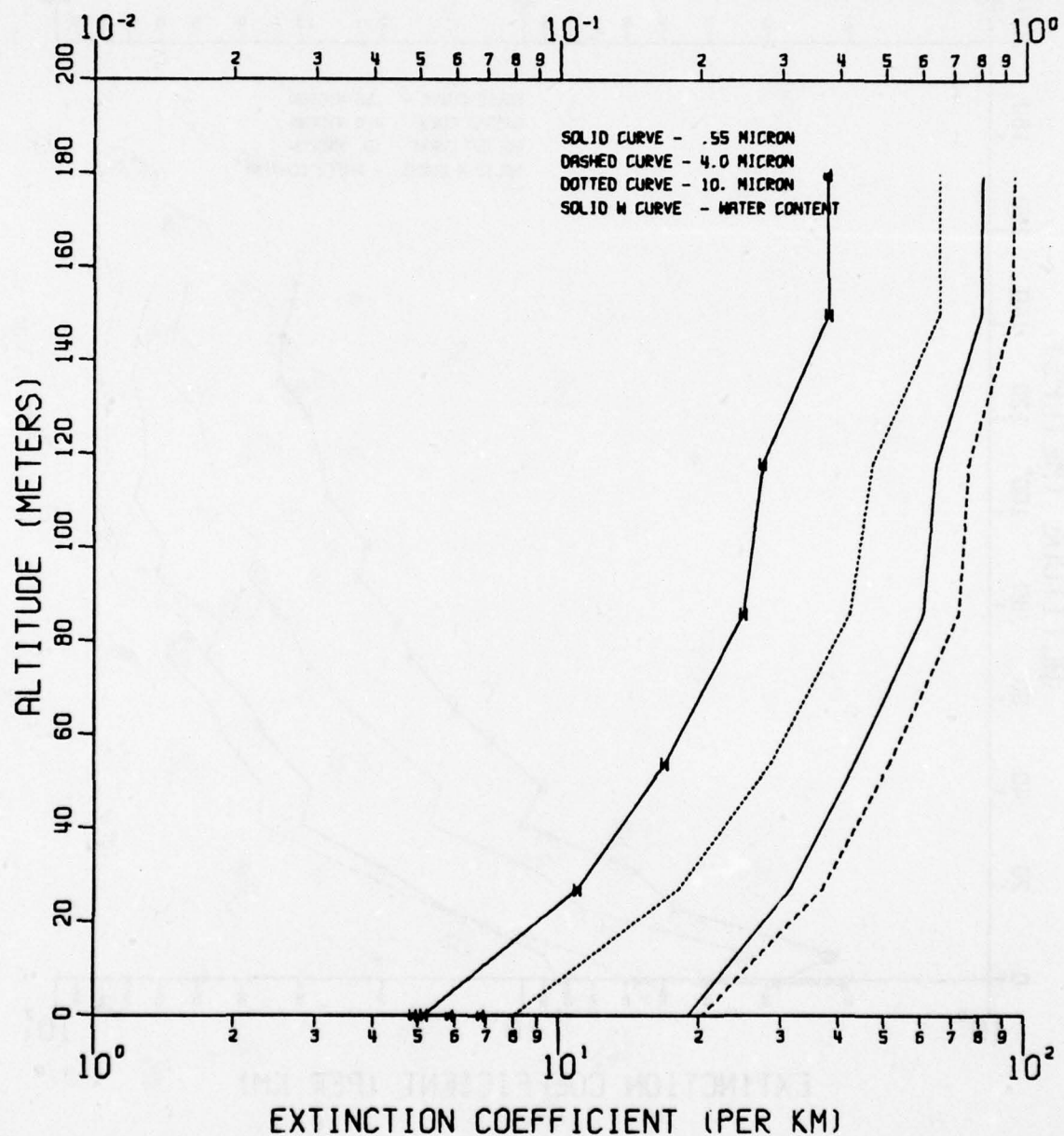


Figure A4-k

DATE = 02/25/76

TIME = 12:25:30 TO 12:31:50

LIQUID WATER CONTENT (GM PER METER**3)

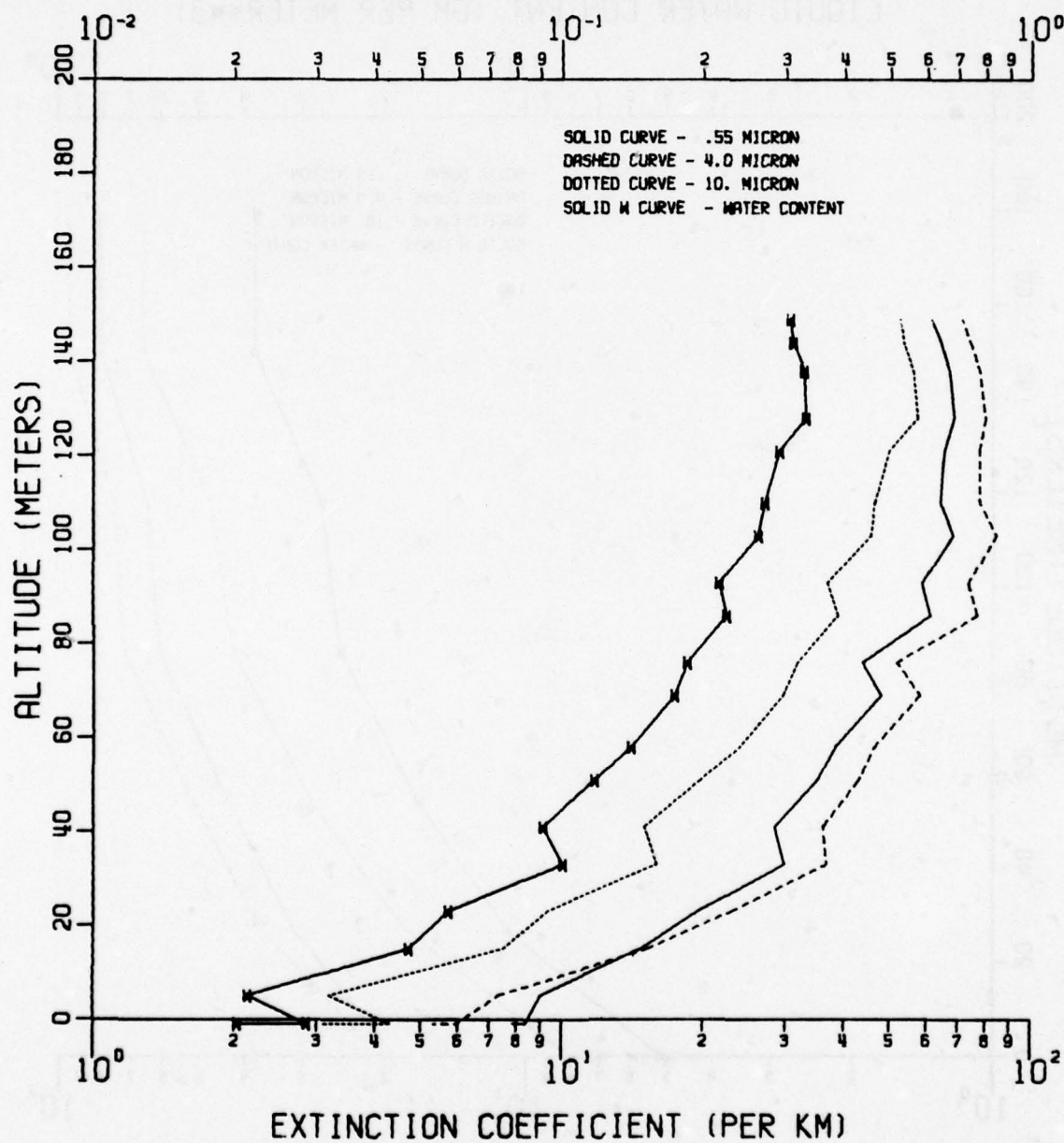


Figure A4-1

DATE = 02/25/76

TIME = 13:20:00 TO 13:26:00

LIQUID WATER CONTENT (GM PER METER**3)

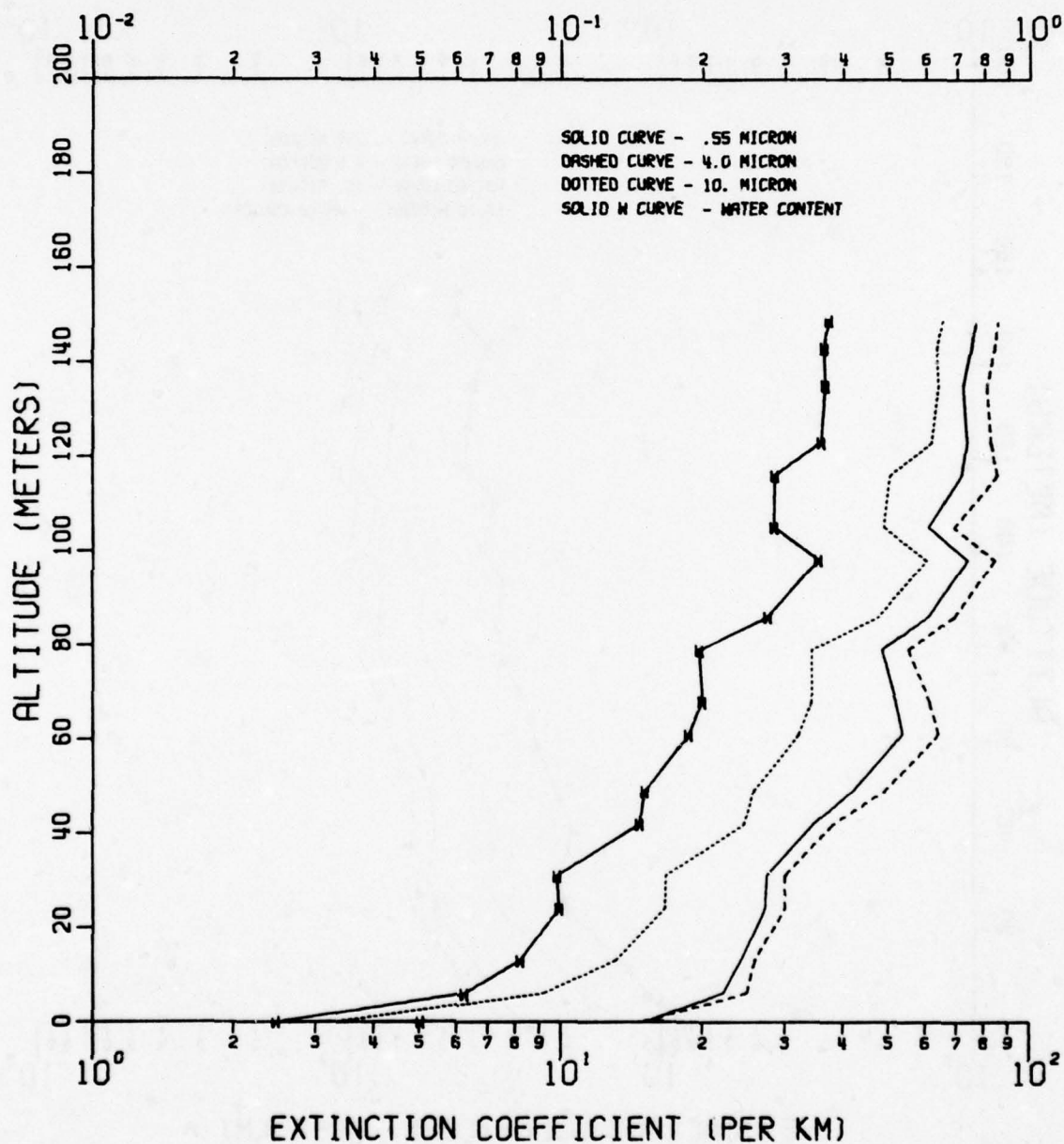


Figure A4-m

A-73

DATE = 02/25/76

TIME = 16:01:40 TO 16:08:20

LIQUID WATER CONTENT (GM PER METER**3)

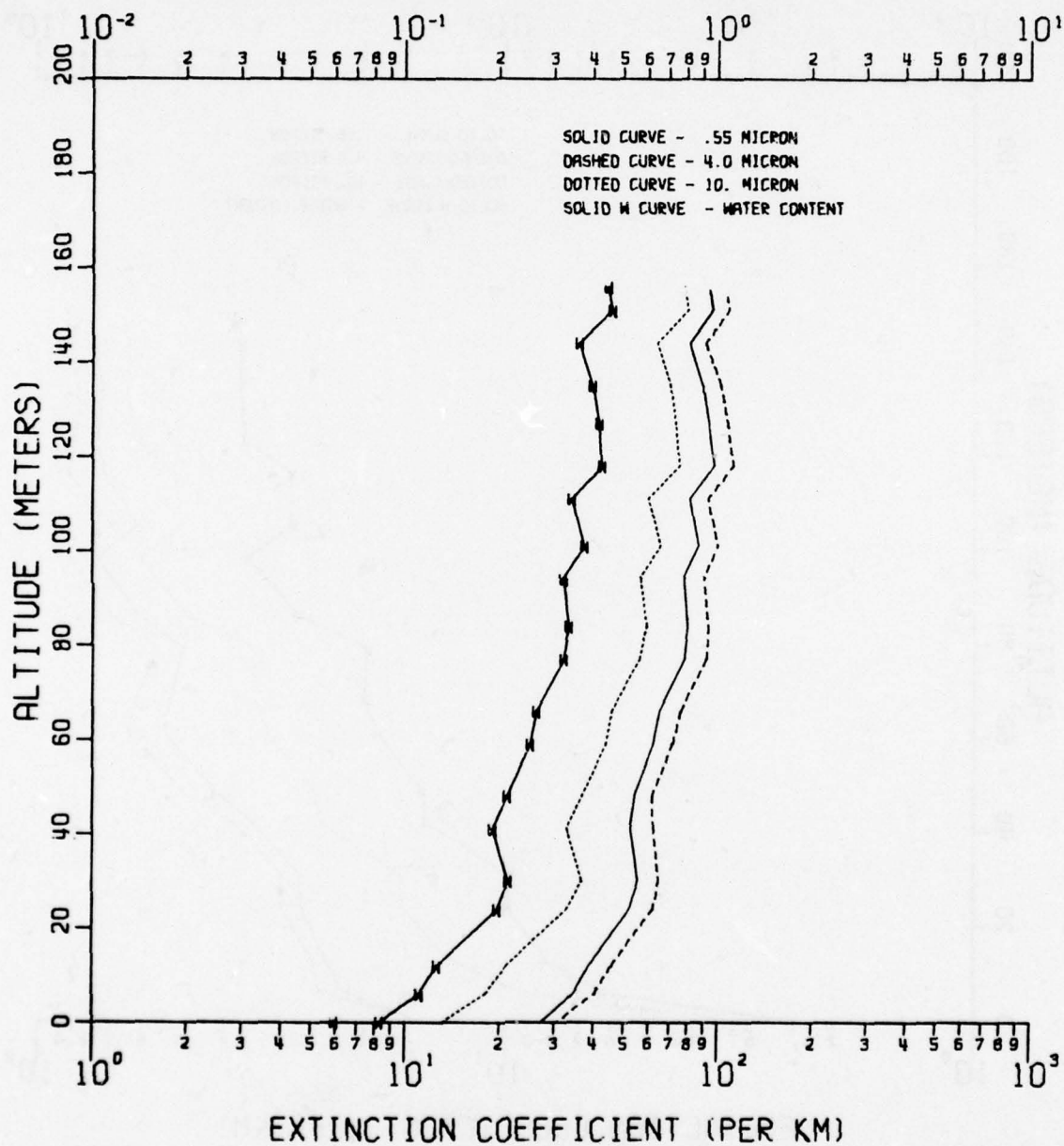


Figure A4-n

DATE = 02/26/76

TIME = 07:32:50 TO 07:40:50

LIQUID WATER CONTENT (GM PER METER**3)

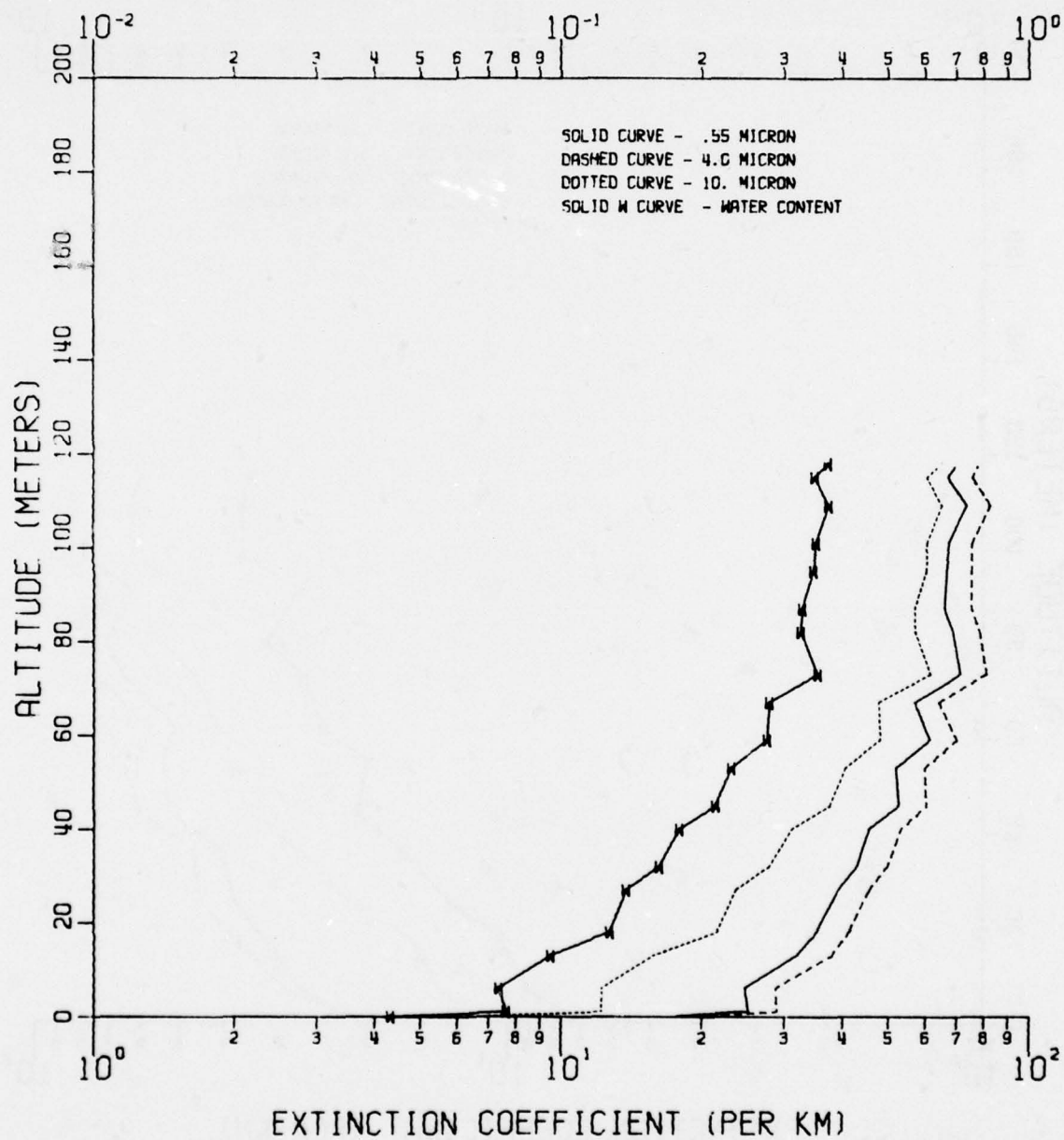


Figure A4-o

DATE = 02/26/76

TIME = 08:24:40 TO 08:29:40

LIQUID WATER CONTENT (GM PER METER**3)

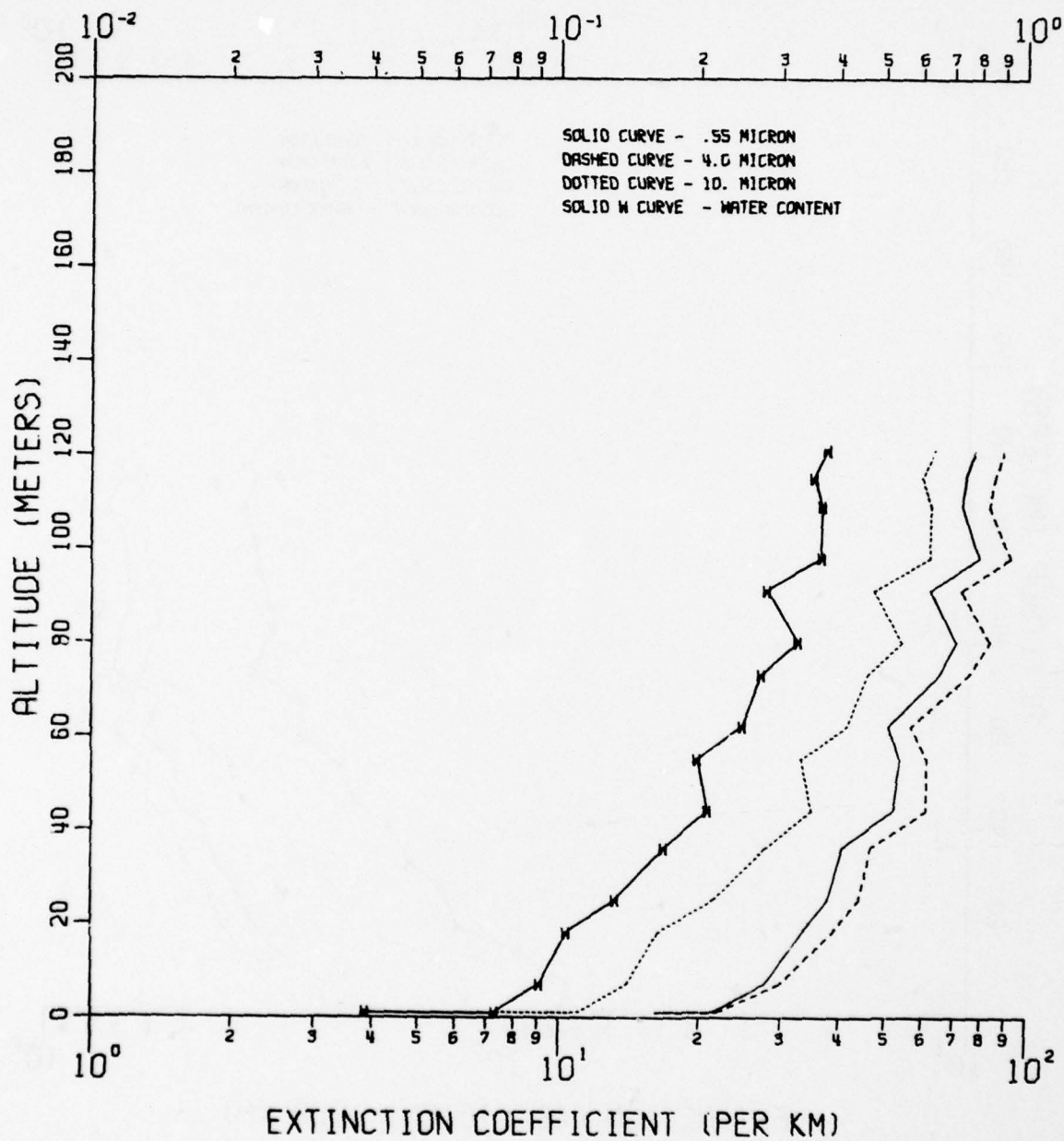


Figure A4-p

DATE = 02/28/76

TIME = 07:55:10 TO 08:01:30

LIQUID WATER CONTENT (GM PER METER**3)

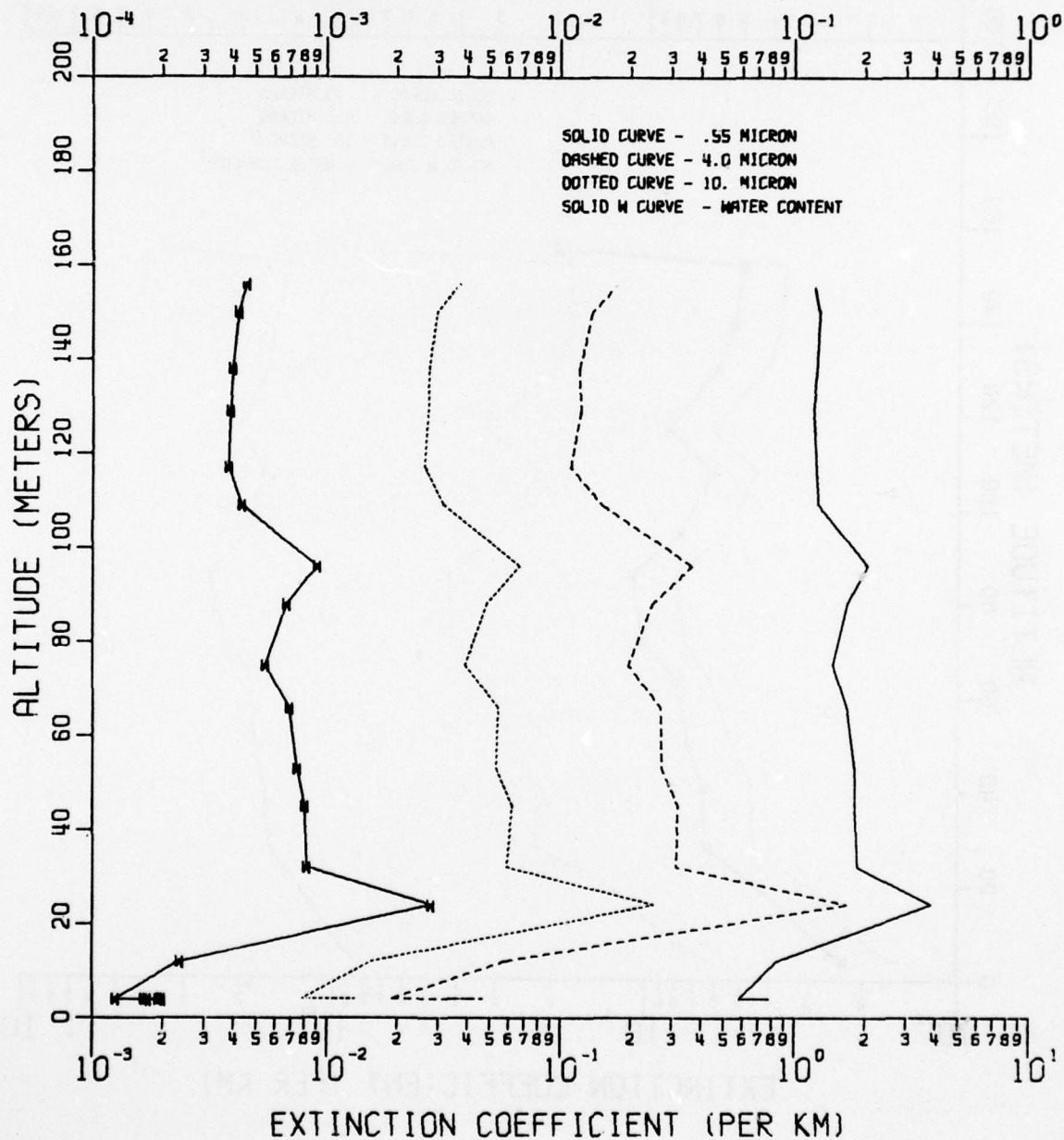


Figure A4-q

DATE = 02/28/76

TIME = 09:30:00 TO 09:35:00

LIQUID WATER CONTENT (GM PER METER**3)

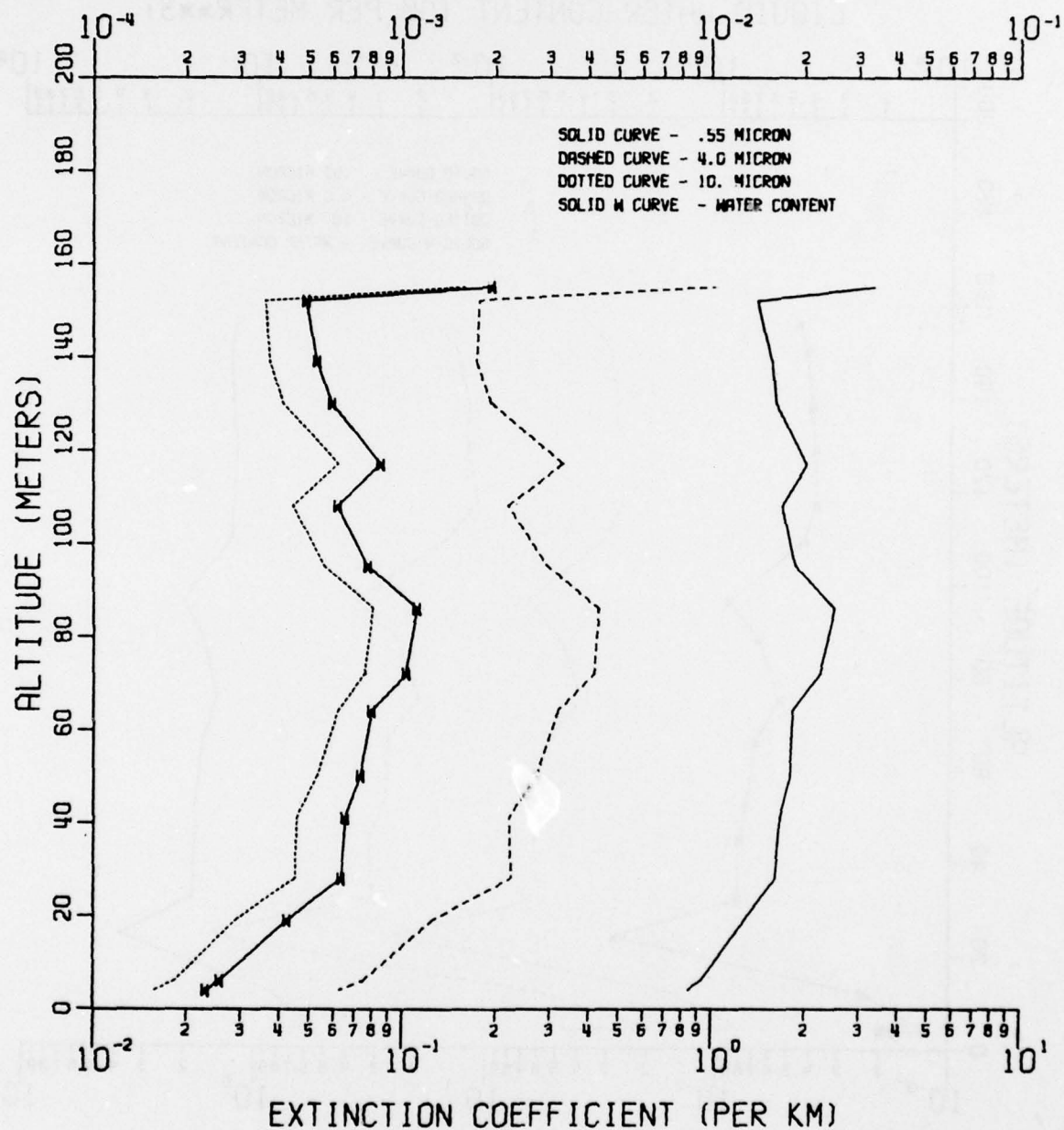


Figure A4-r

DATE = 02/28/76

TIME = 10:17:00 TO 10:22:20

LIQUID WATER CONTENT (GM PER METER**3)

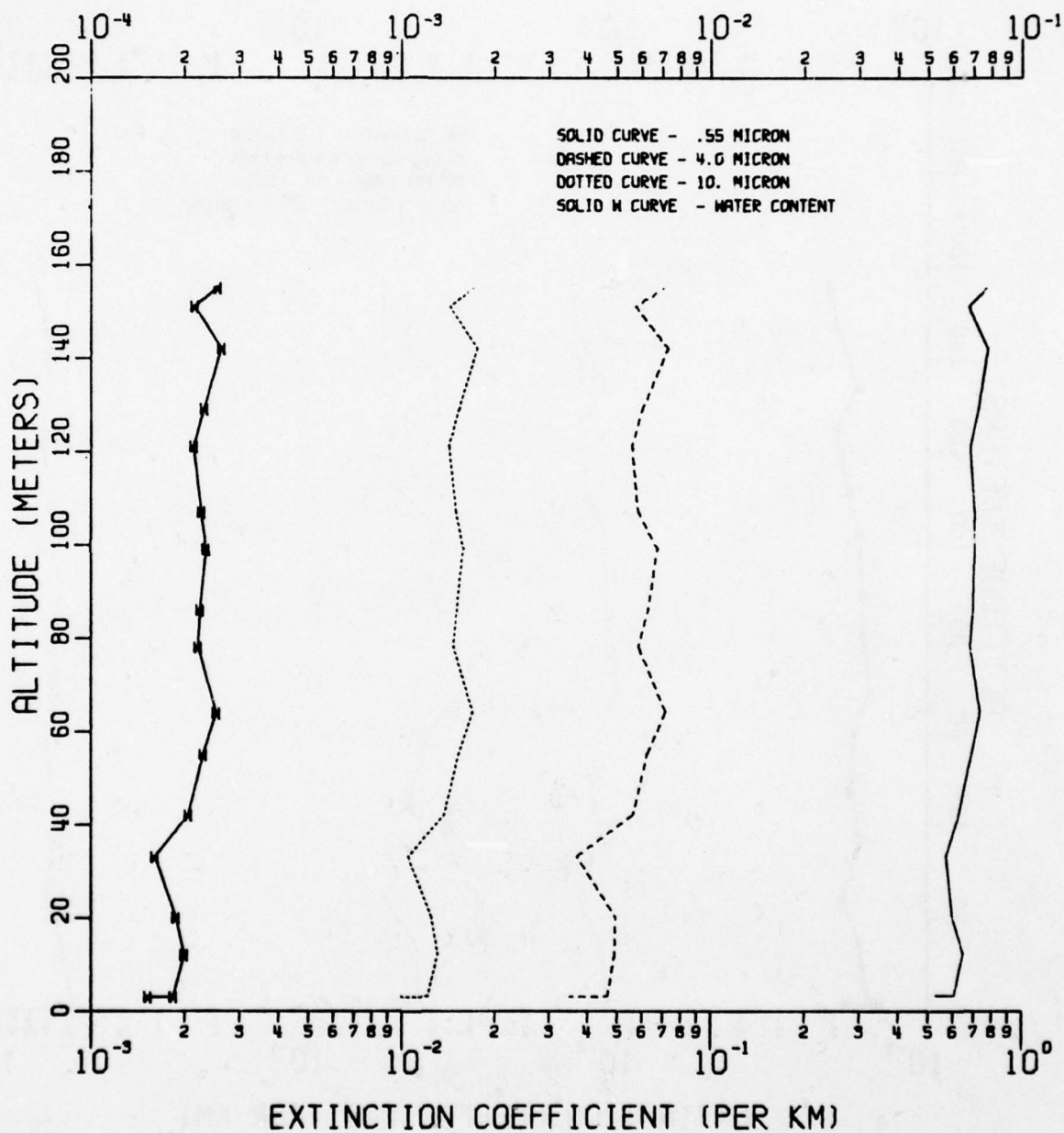


Figure A4-s

DATE = 02/28/76

TIME = 11:07:10 TO 11:12:50

LIQUID WATER CONTENT (GM PER METER**3)

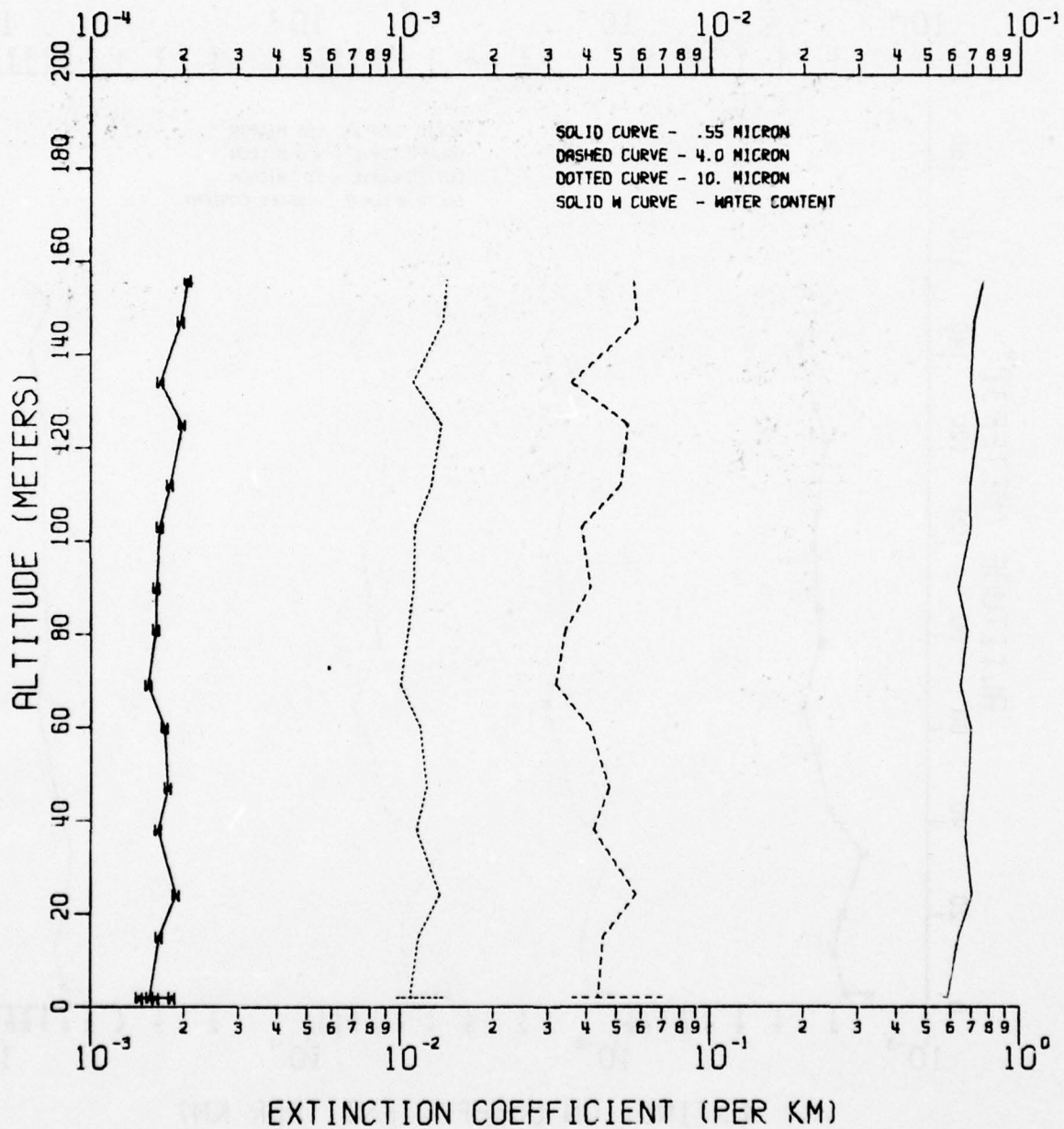


Figure A4-t

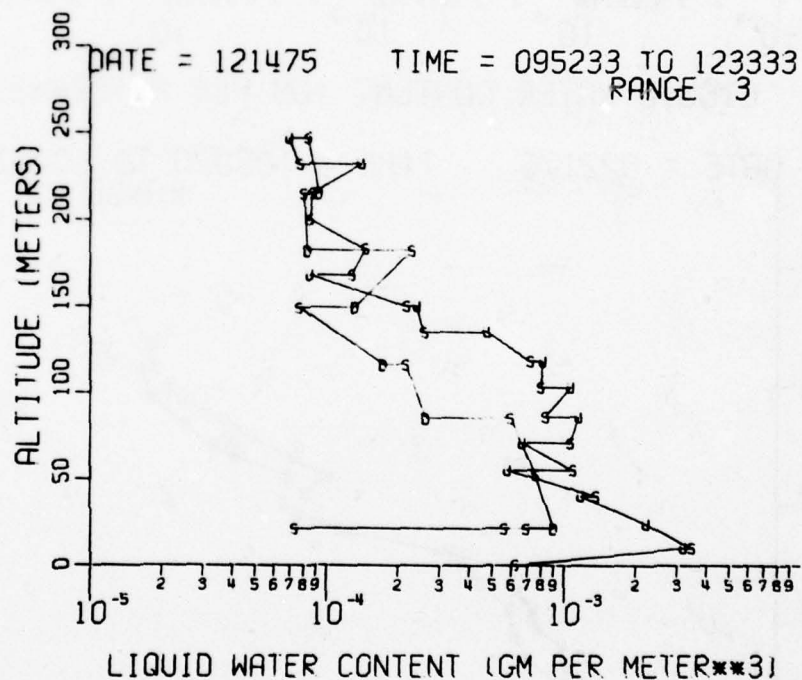
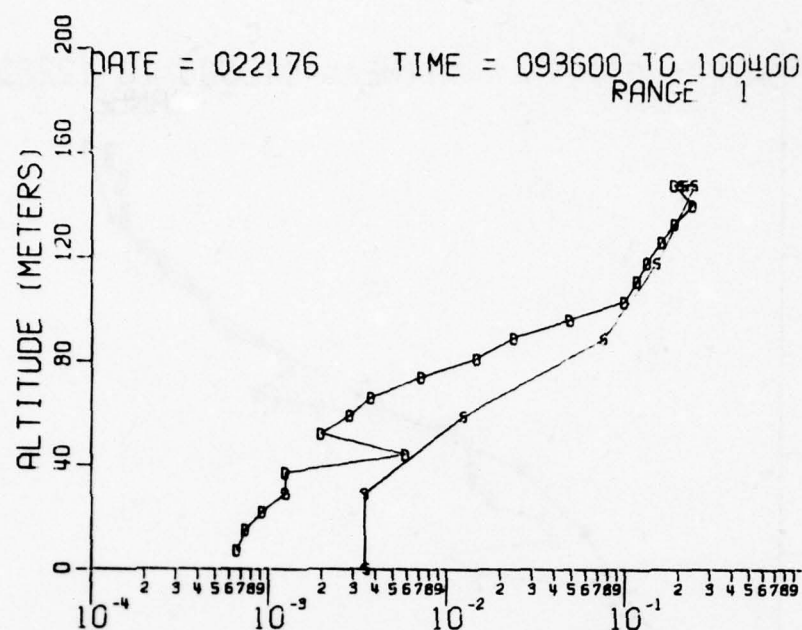


Figure A5-a. Profiles of liquid water content inferred from particle size distribution measurements (shown in Fig. A1) made with a balloon-borne aerosol counter. Both upleg and downleg portions of each traverse are presented. The u, s, and d symbols indicate the aerosol counter was moving upward, was stationary, or was moving downward during the measurement period. The date, local time (hours, minutes, seconds) and time interval for the measurements are given, along with the size range setting (see Fig. 4) of the aerosol counter.

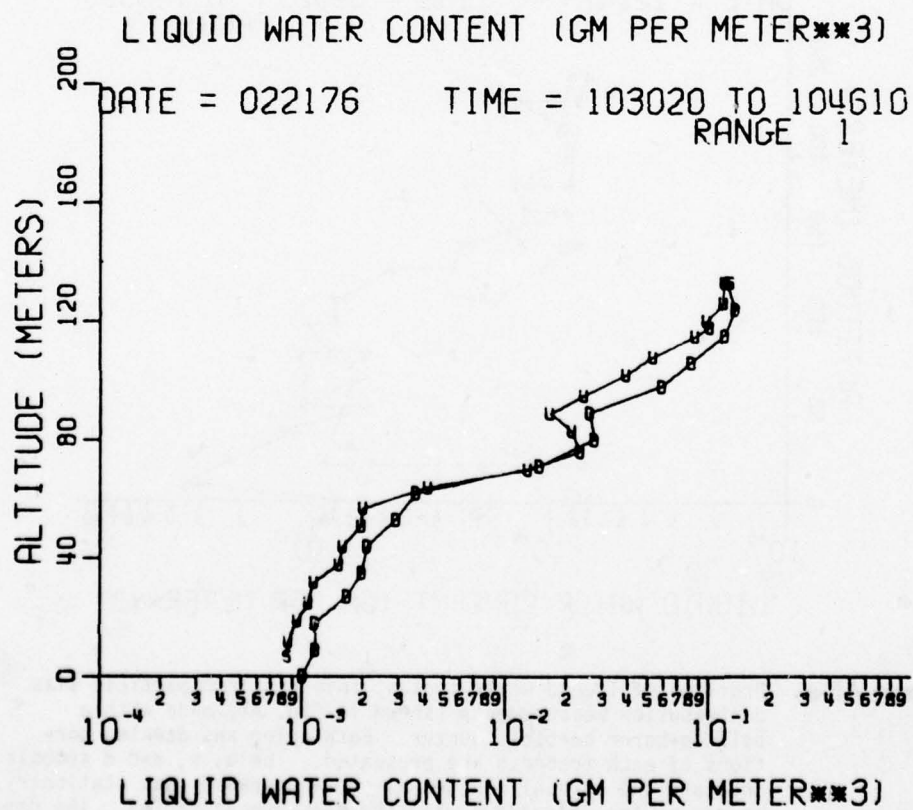
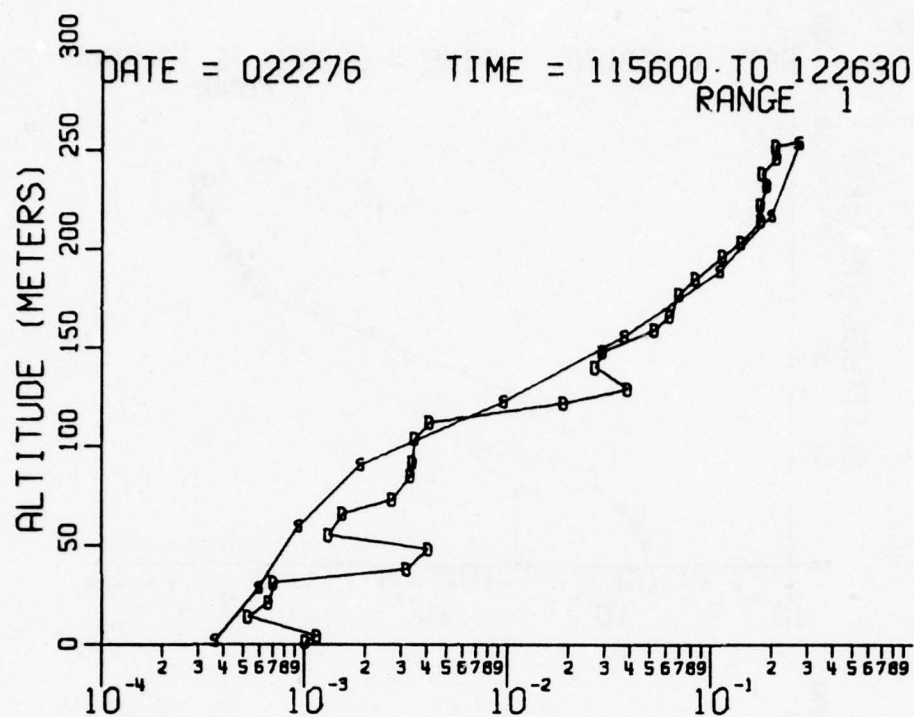


Figure A5-b

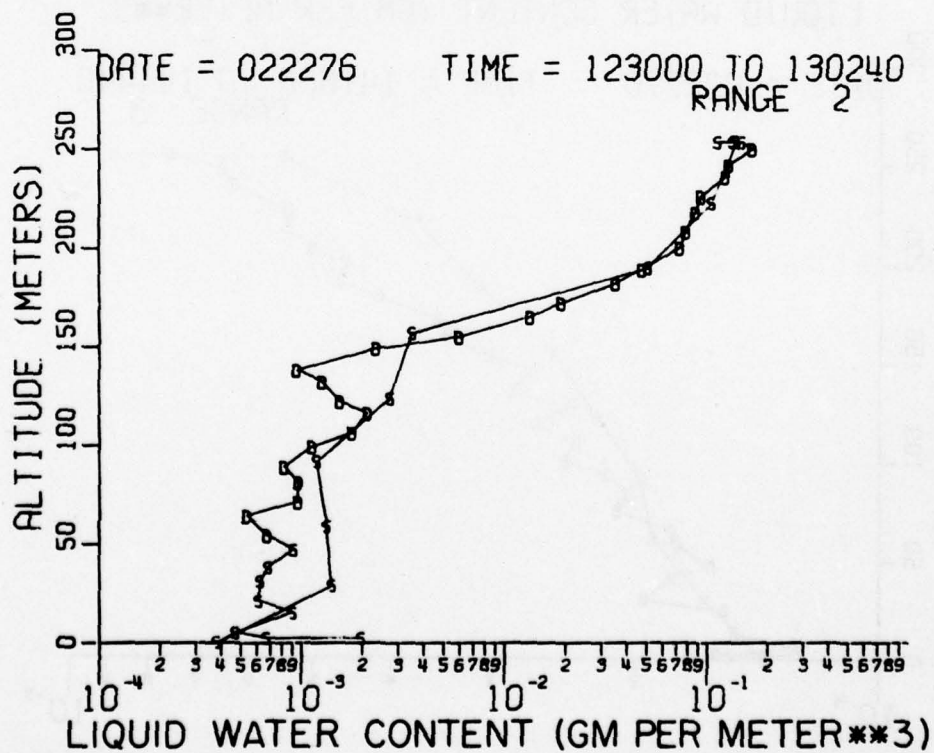
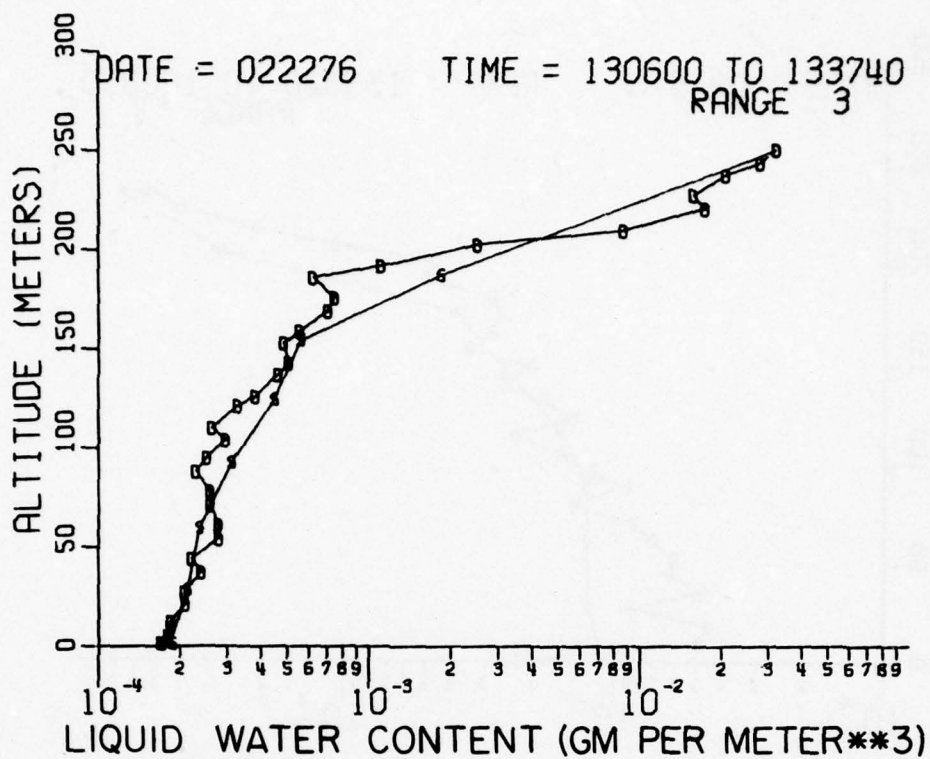


Figure A5-c

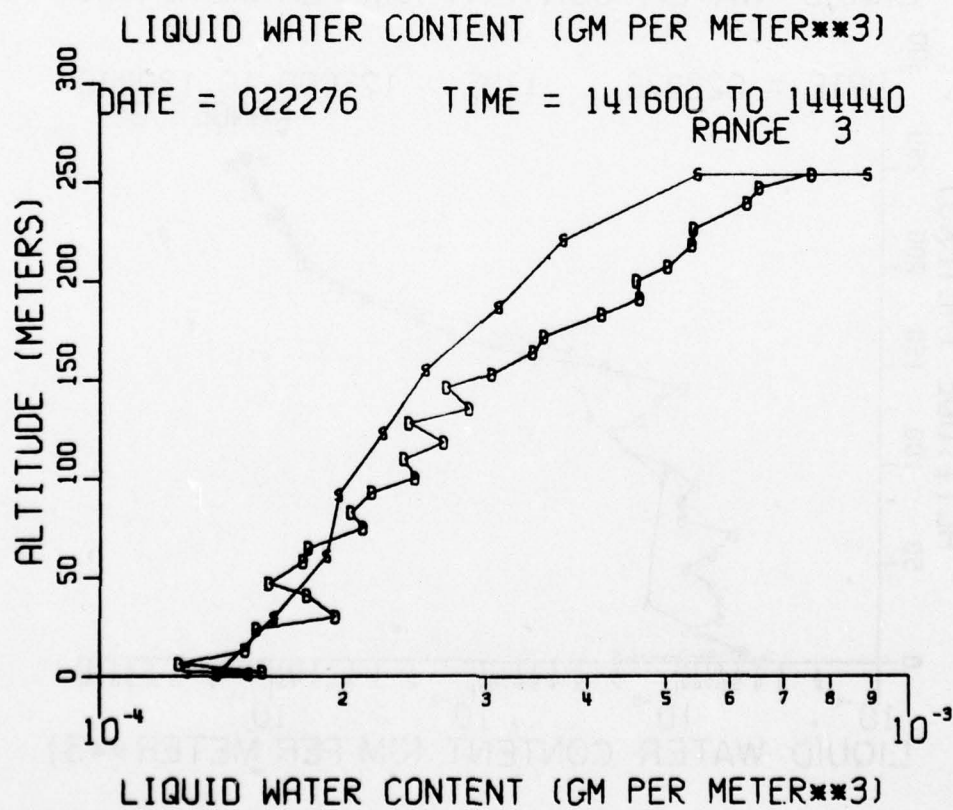
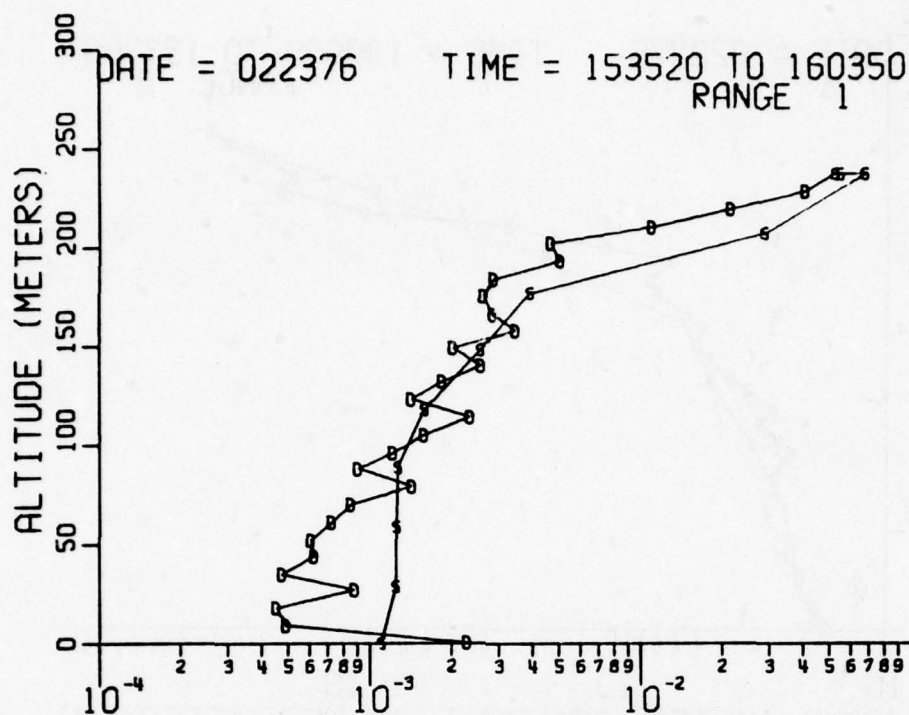


Figure A5-d

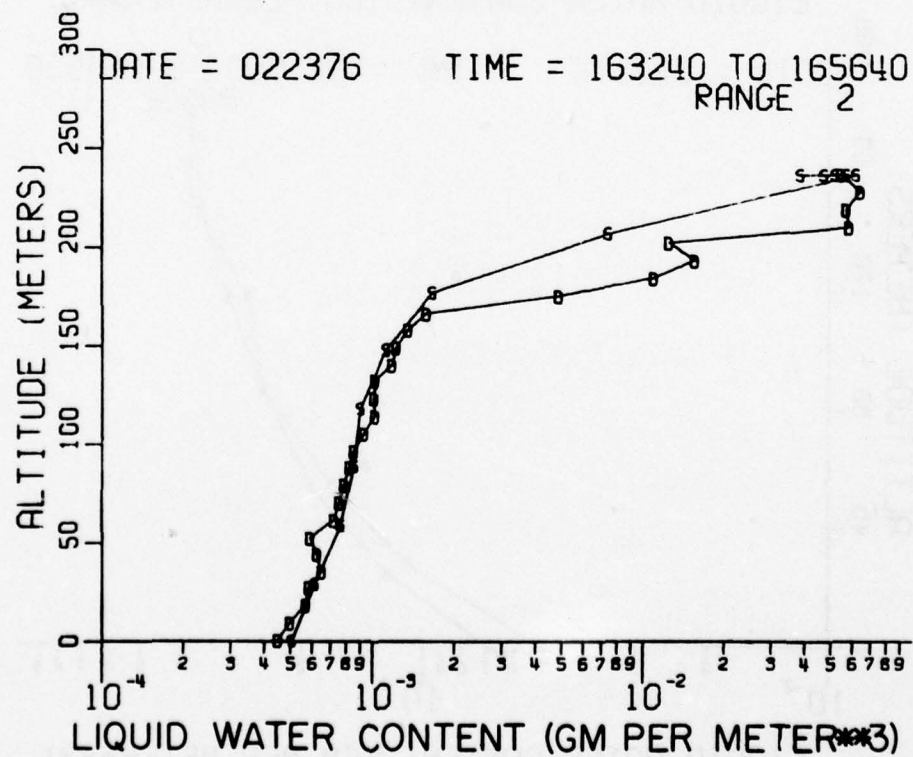
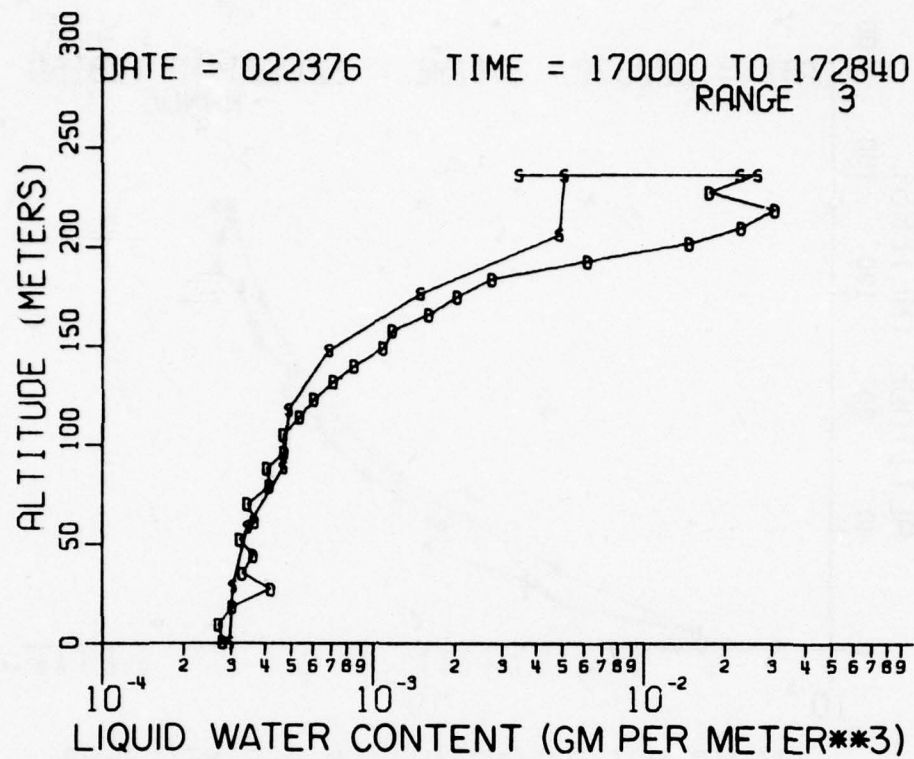


Figure A5-e

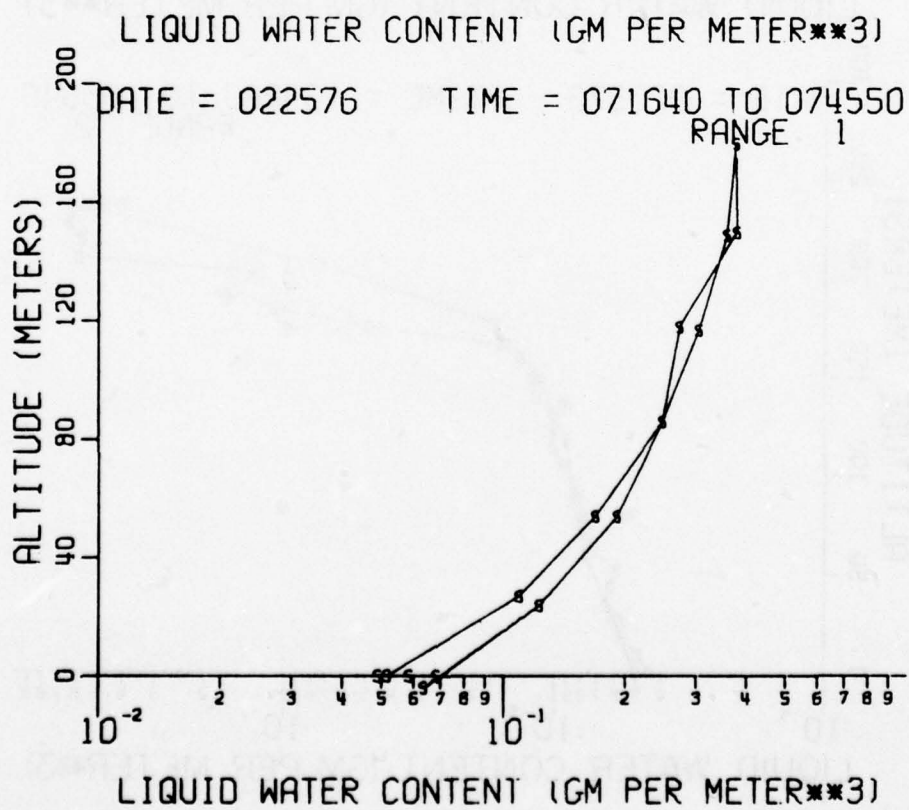
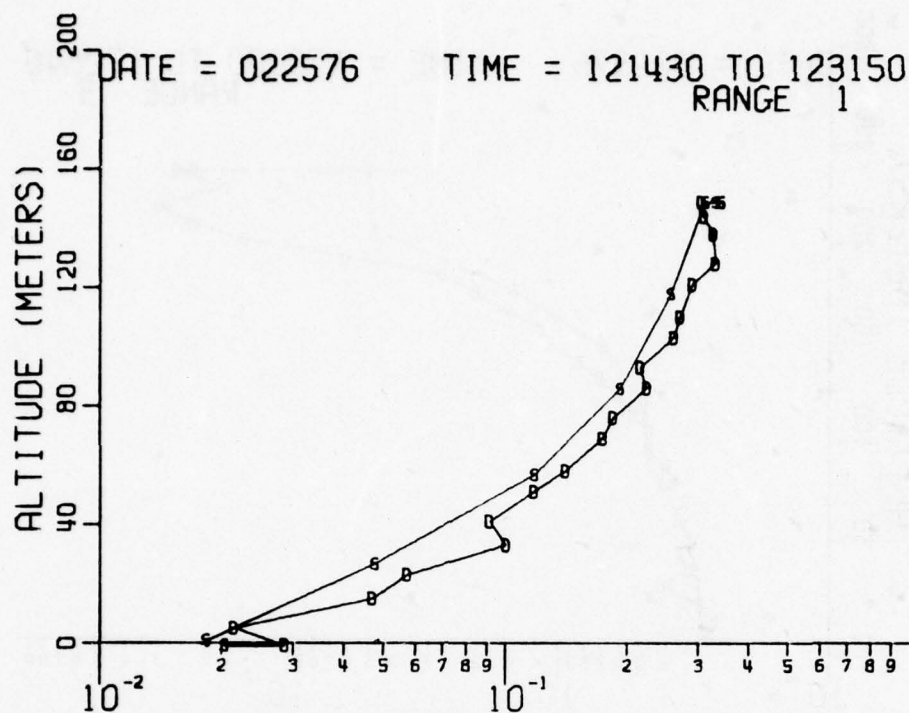


Figure A5-f

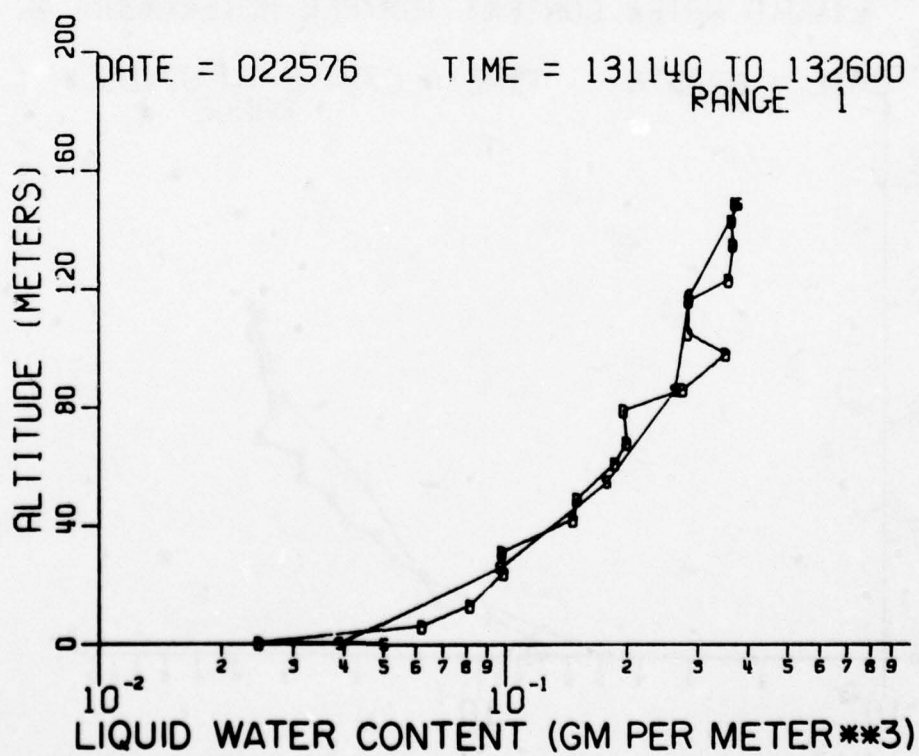
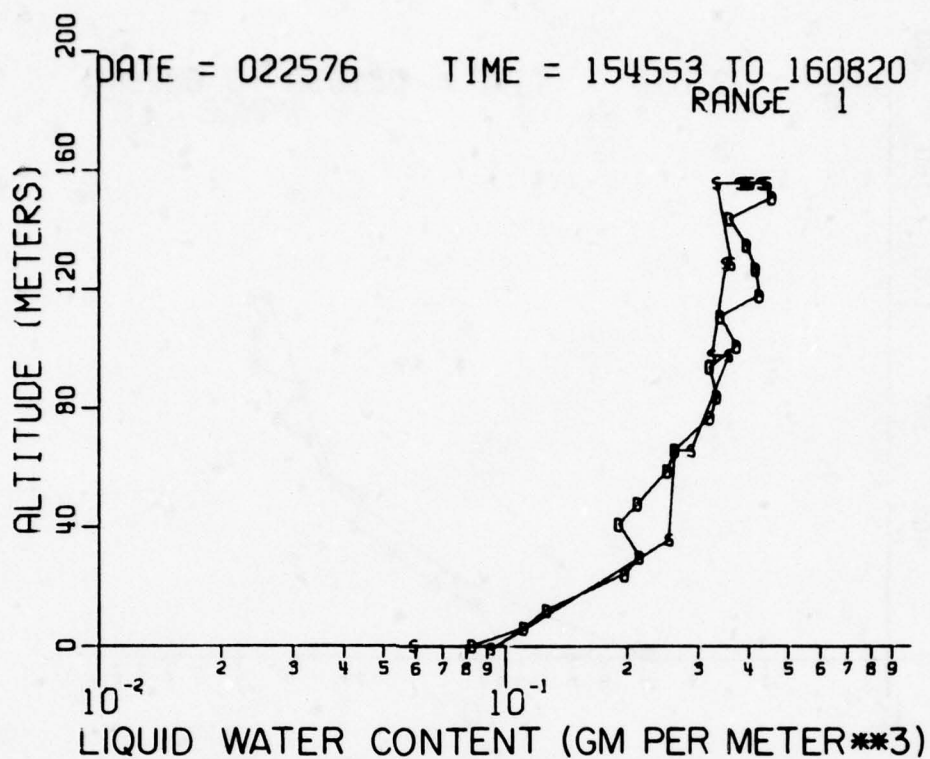


Figure A5-g

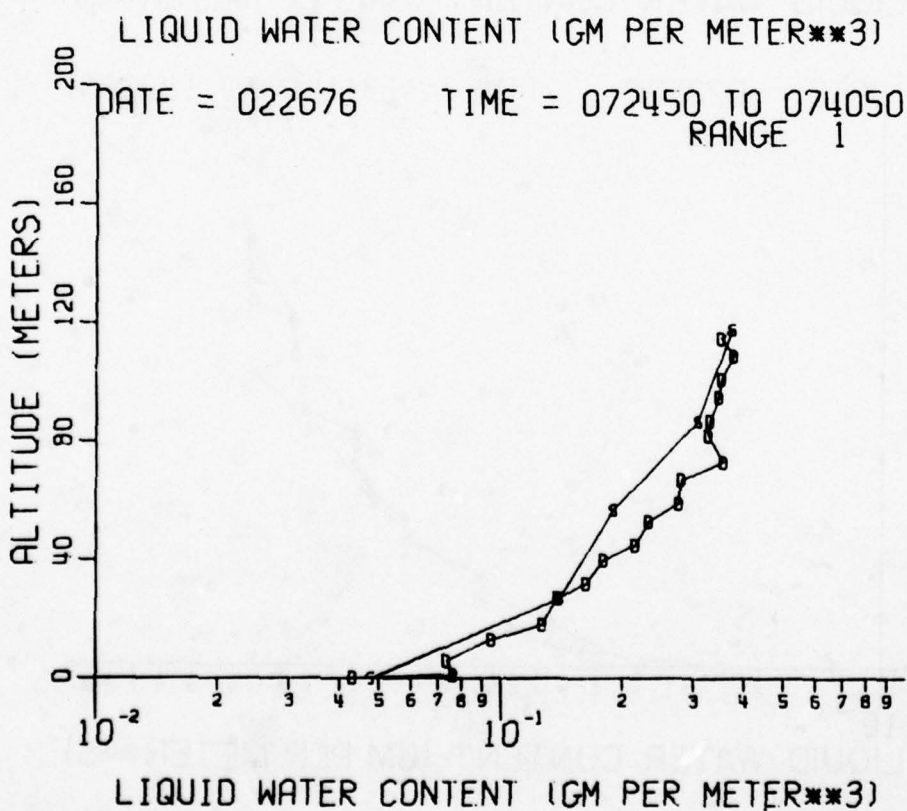
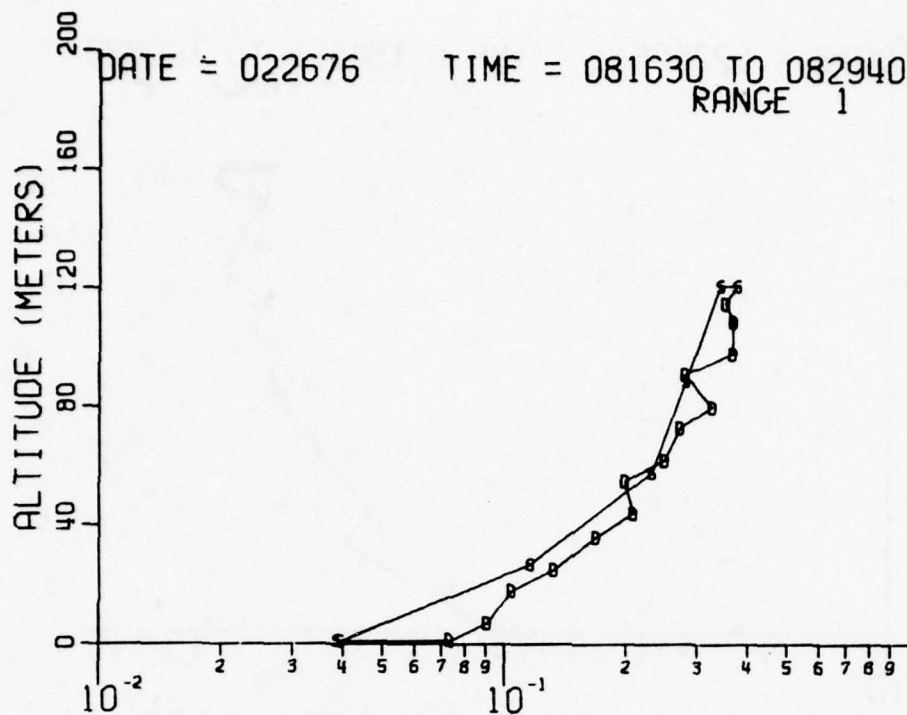


Figure A5-h

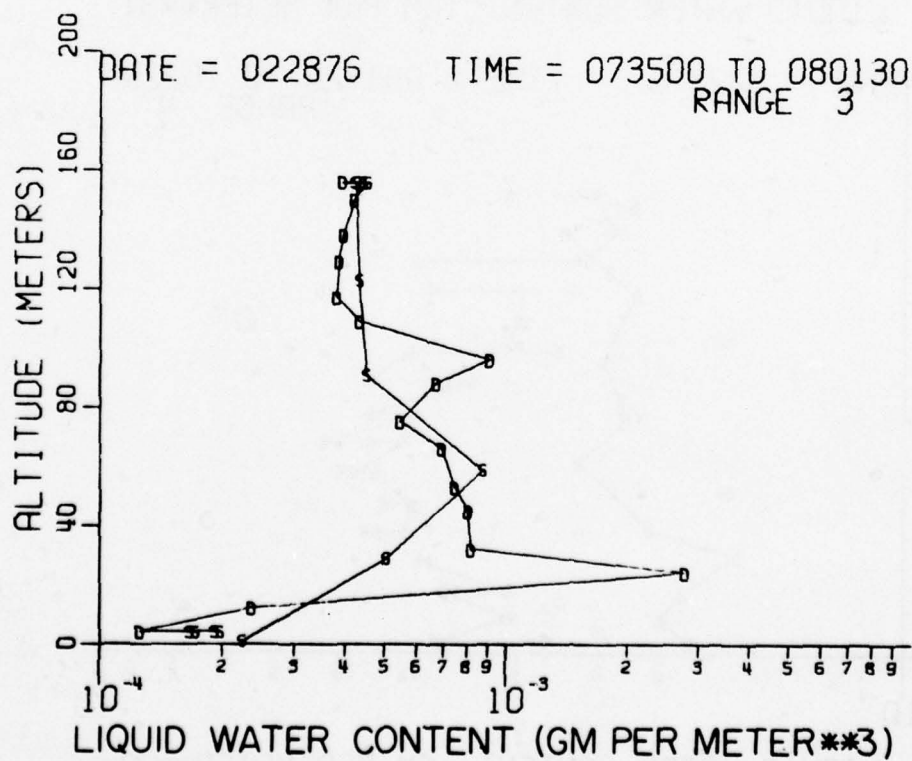
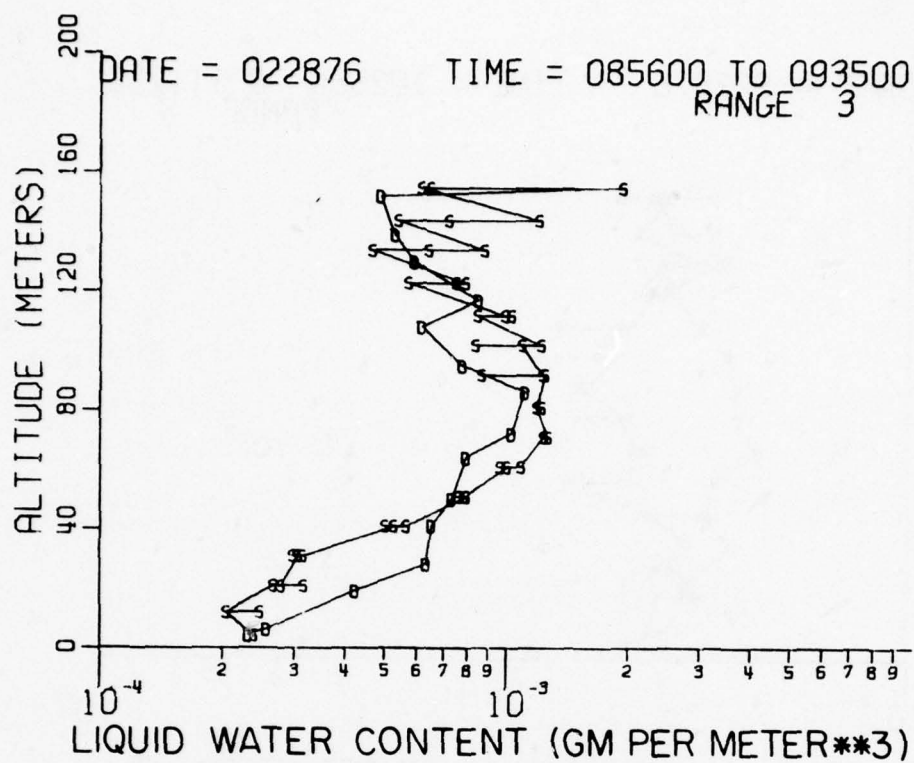


Figure A5-i

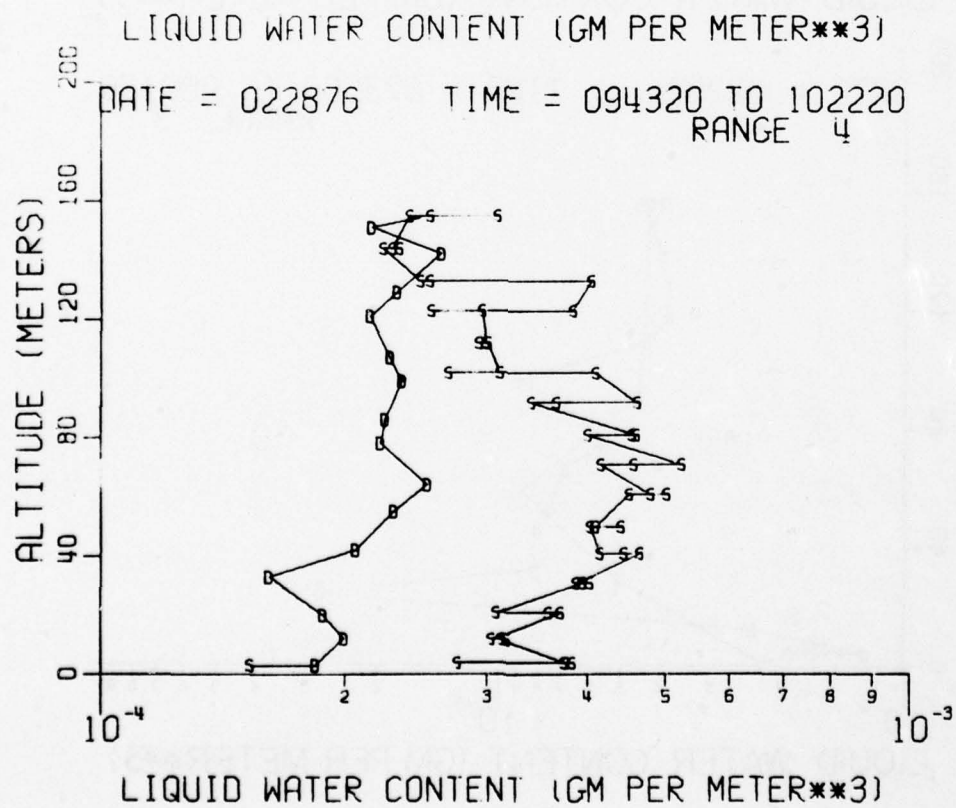
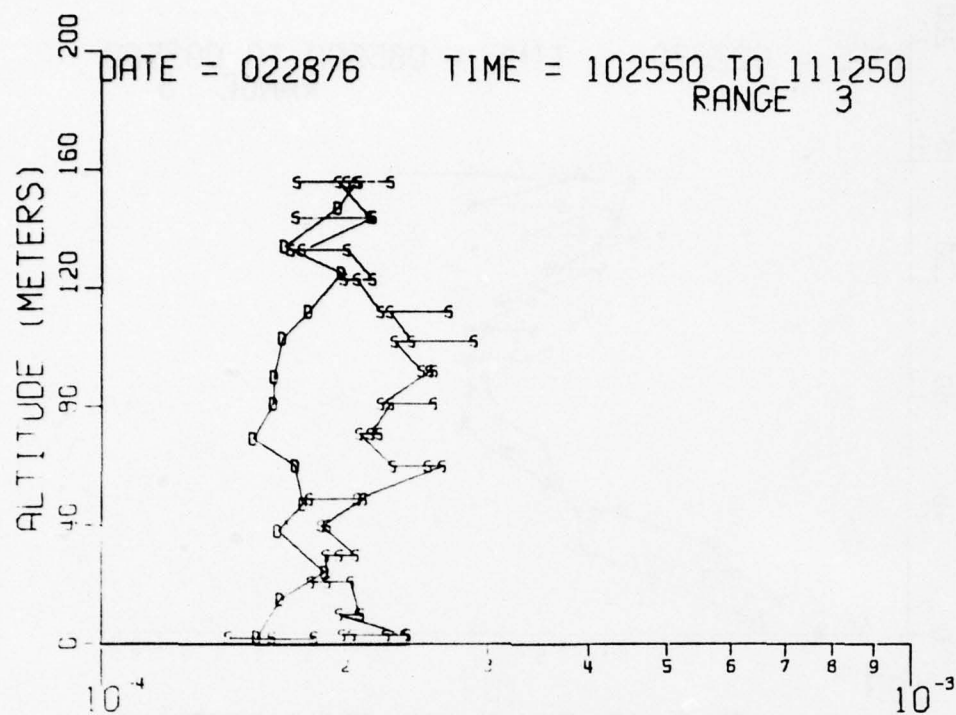


Figure A5-j

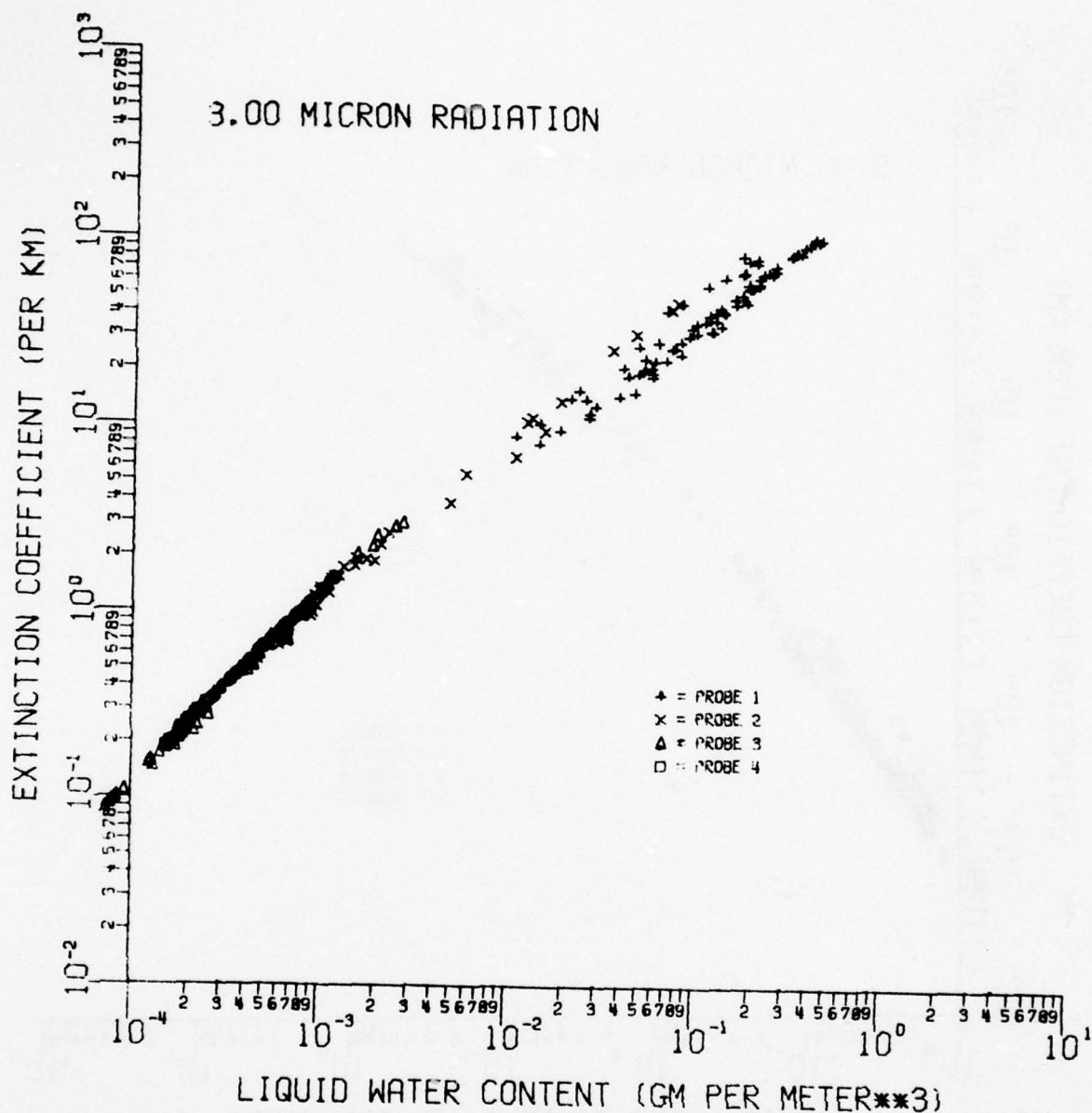


Figure A6-a. Values of the Mie theory calculated particulate extinction at $3\mu\text{m}$, $5\mu\text{m}$, $8\mu\text{m}$, $11\mu\text{m}$, and $12\mu\text{m}$ wavelengths versus liquid water content for a variety of particle size distributions of atmospheric fog and haze. The probe range setting 1-4 indicates the particle counter used to make the size distribution measurements on which this data is based was set on different size ranges of sensitivity (as per Fig. 4).

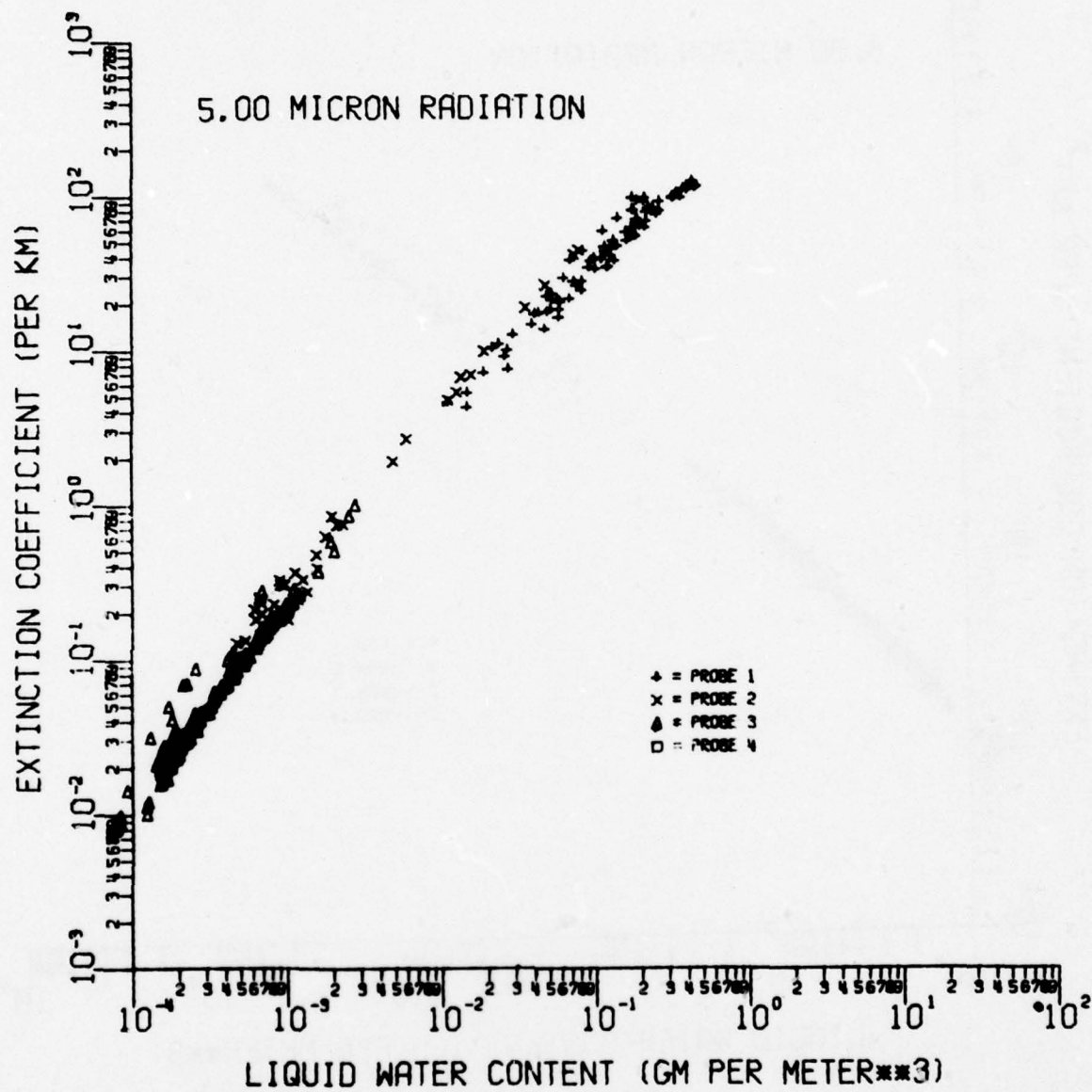


Figure A6-b

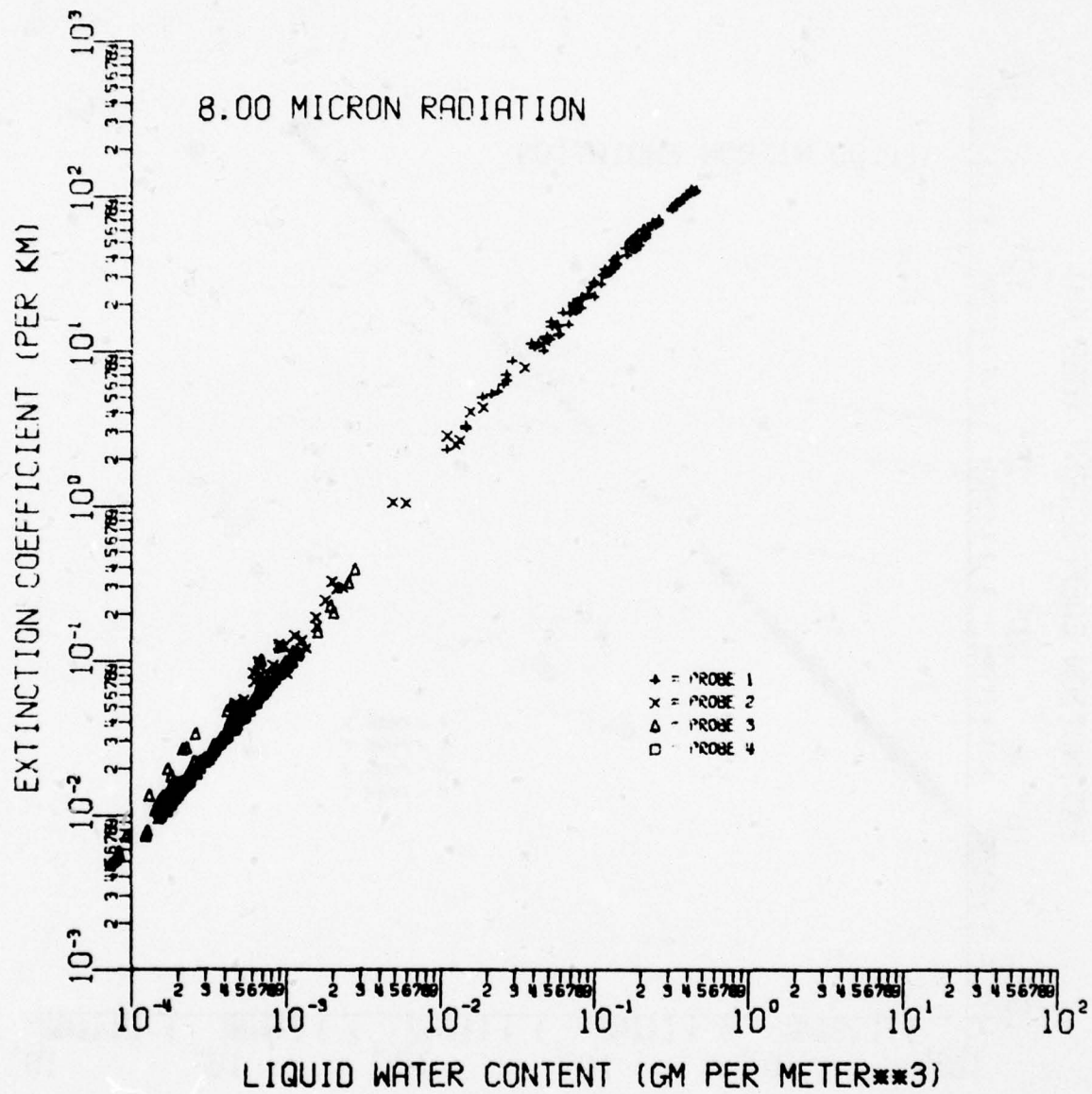


Figure A6-c

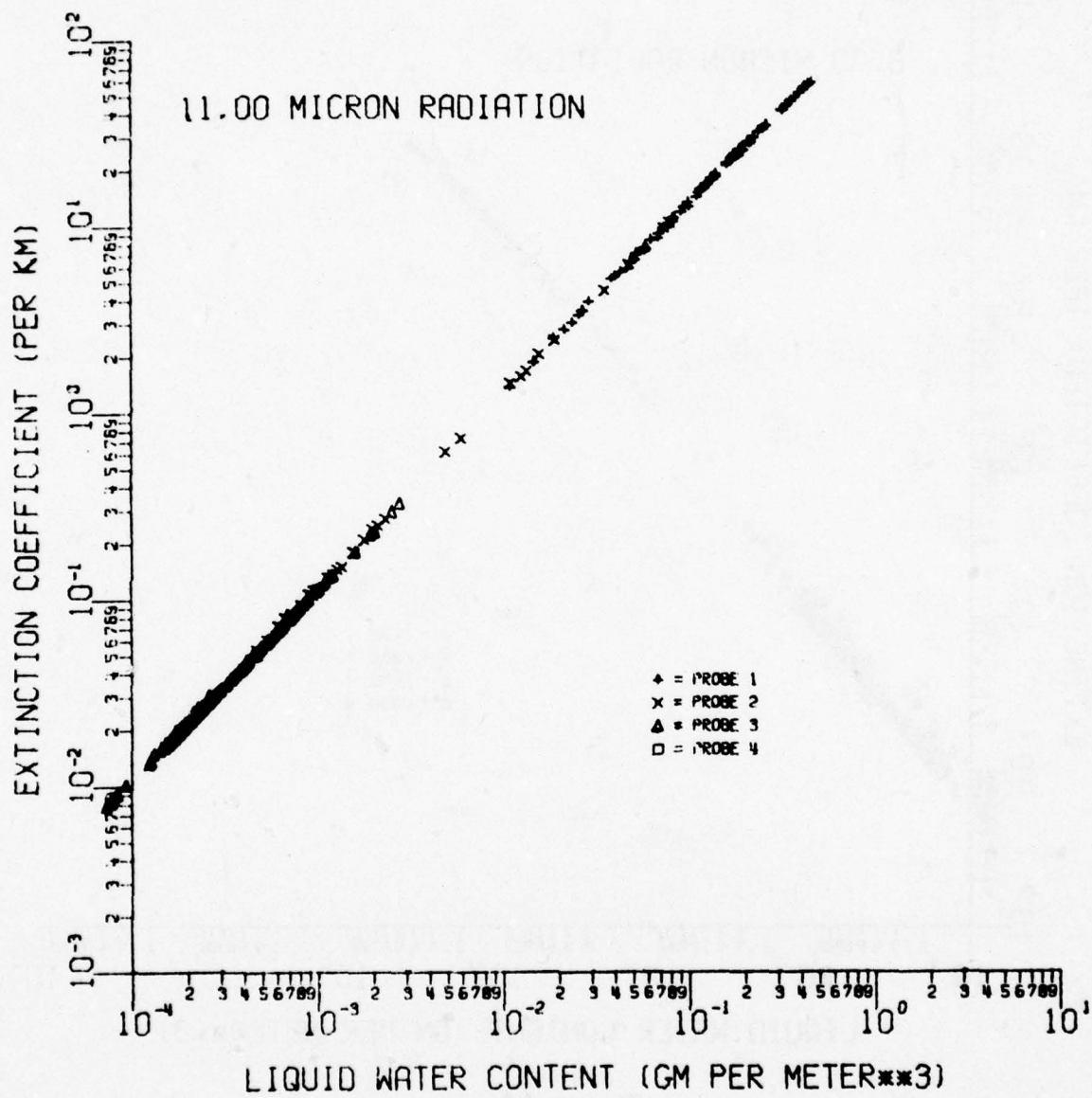


Figure A6-d

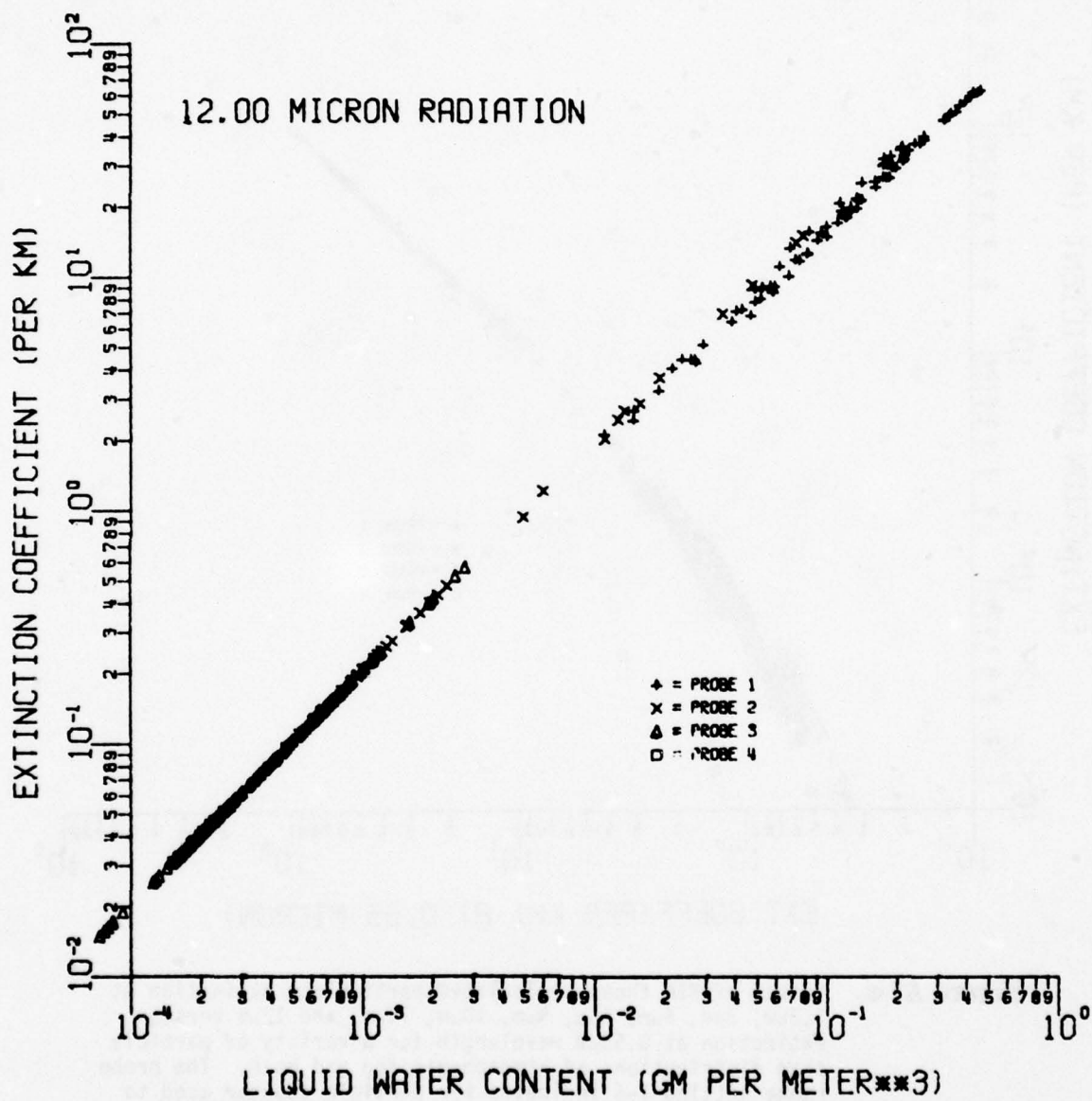


Figure A6-e

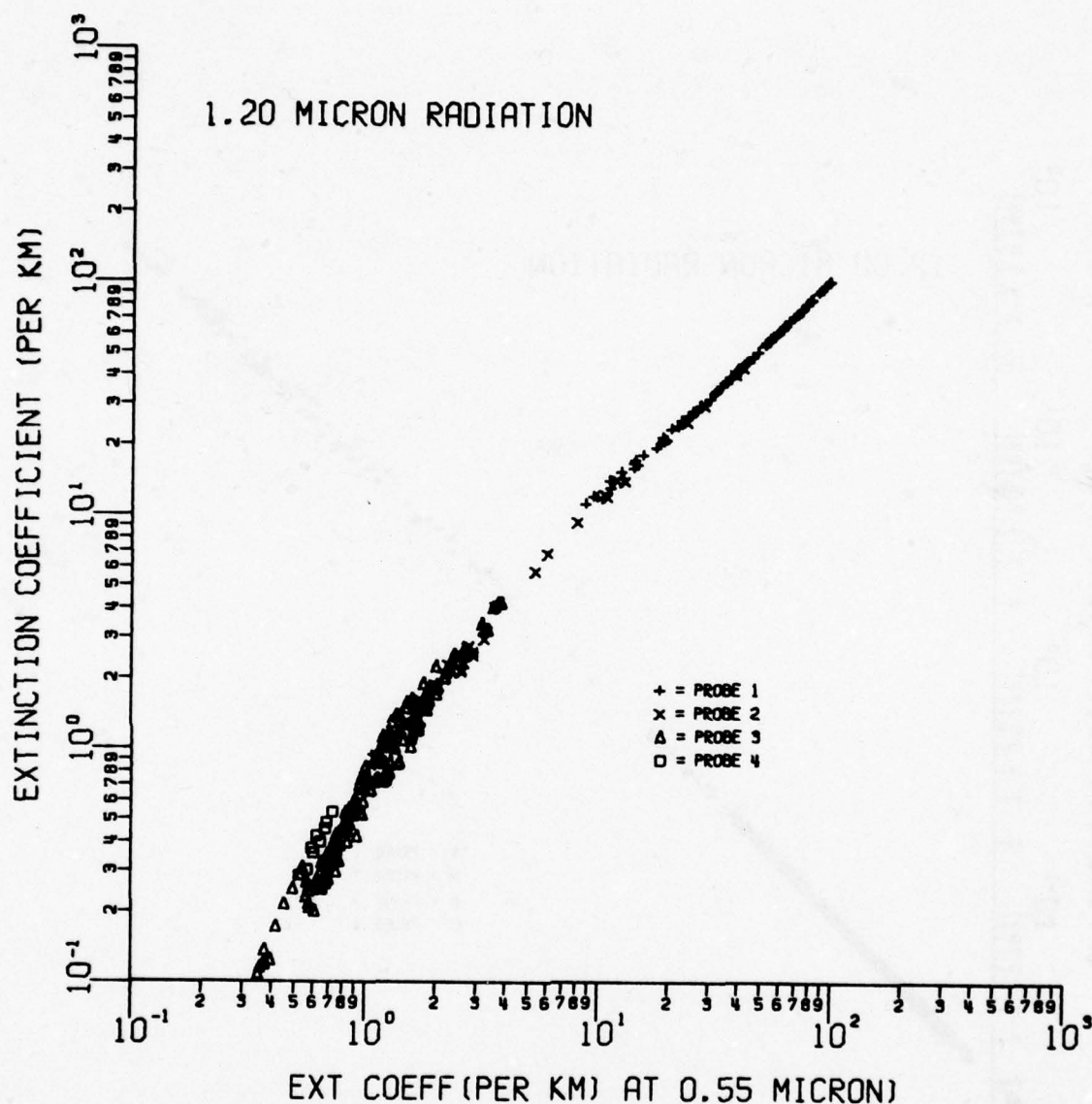


Figure A7-a. Values of Mie theory calculated particulate extinction at 1.2 μ m, 3 μ m, 4 μ m, 5 μ m, 8 μ m, 10 μ m, 11 μ m, and 12 μ m versus extinction at 0.55 μ m wavelength for a variety of particle size distributions of atmospheric fog and haze. The probe range setting 1-4 indicates the particle counter used to make the size distribution measurements on which this data is based was set on different size ranges of sensitivity (as per Fig. 4).

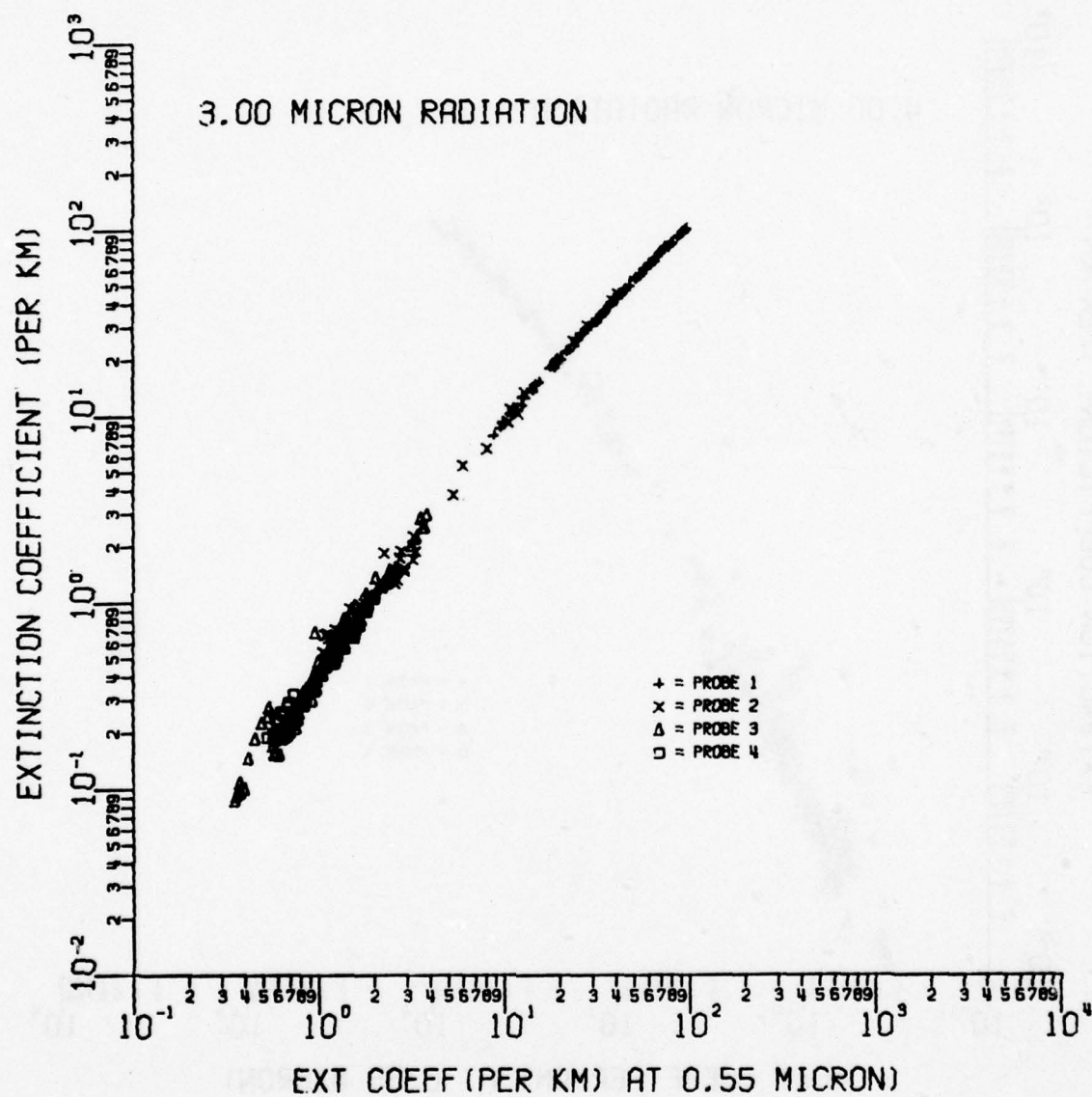


Figure A7-b

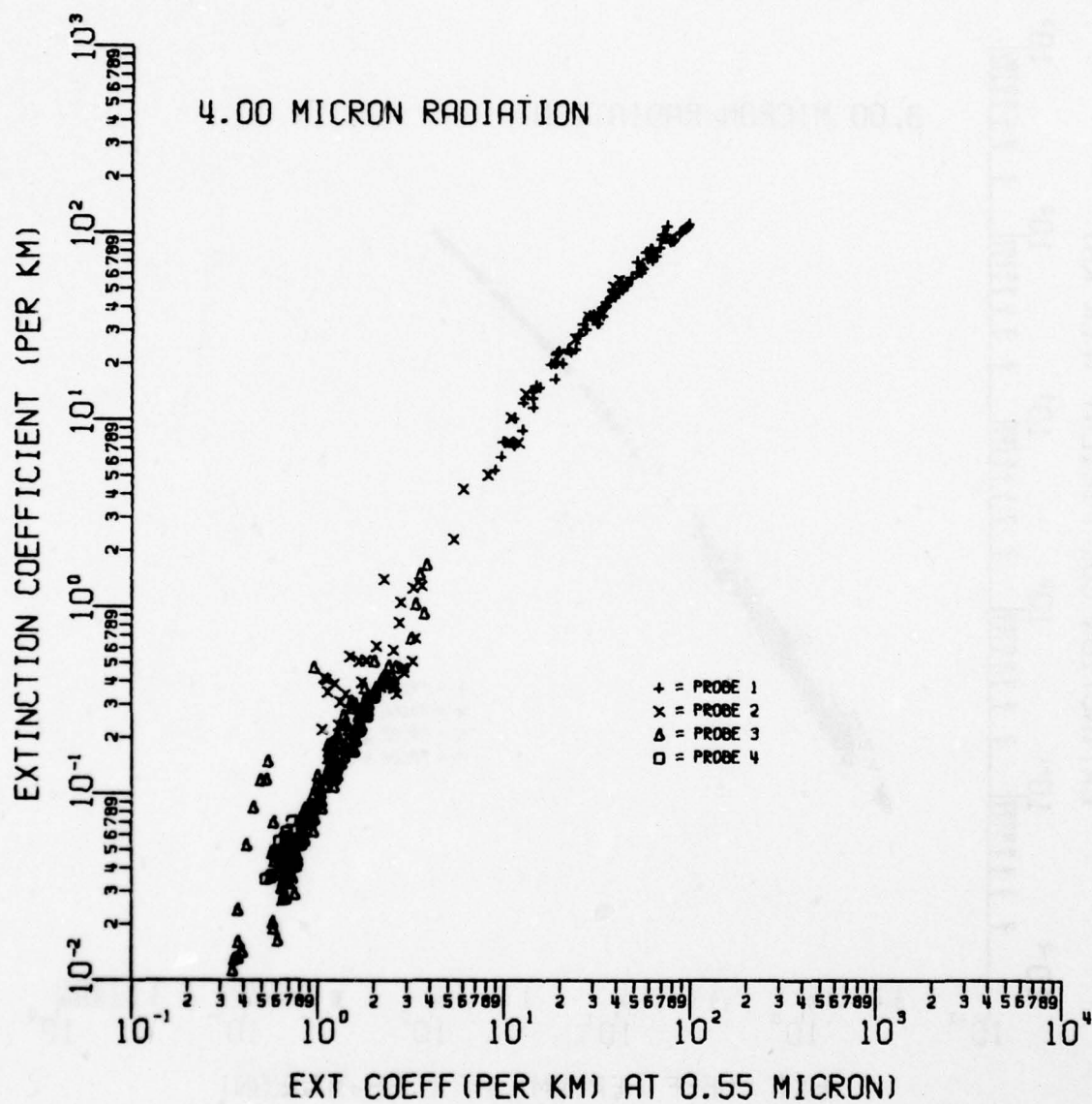


Figure A7-c

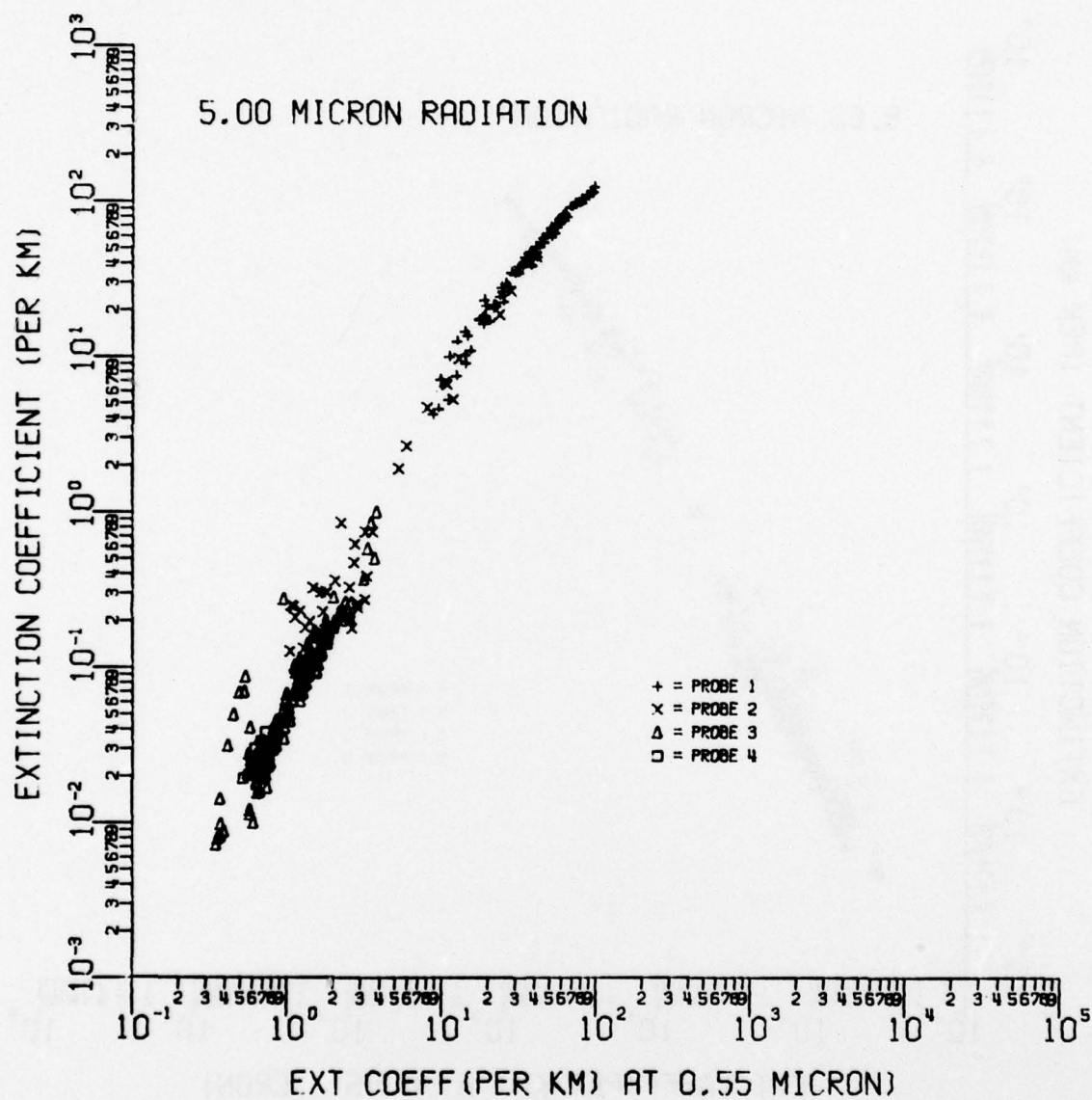


Figure A7-d

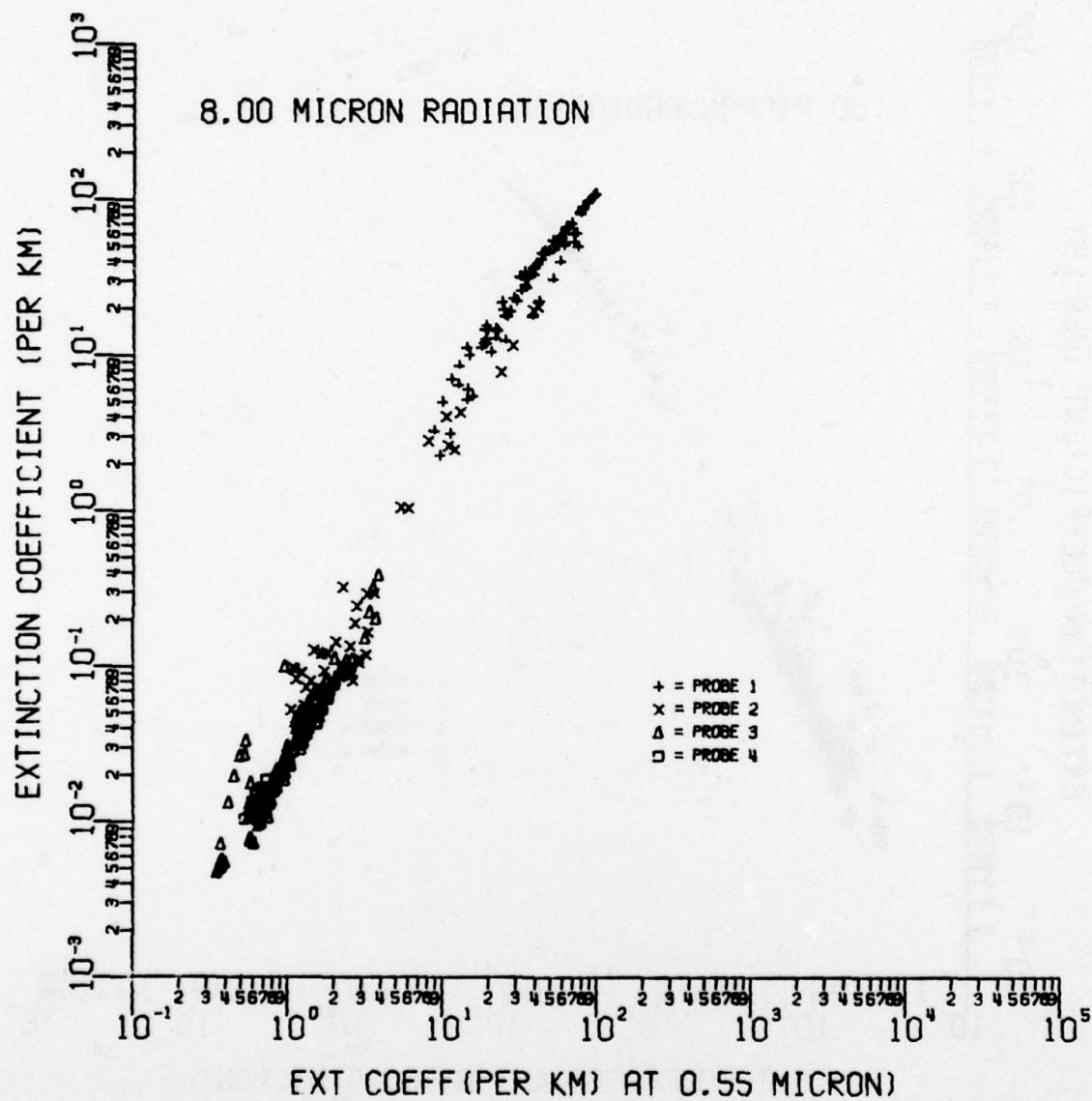


Figure A7-e

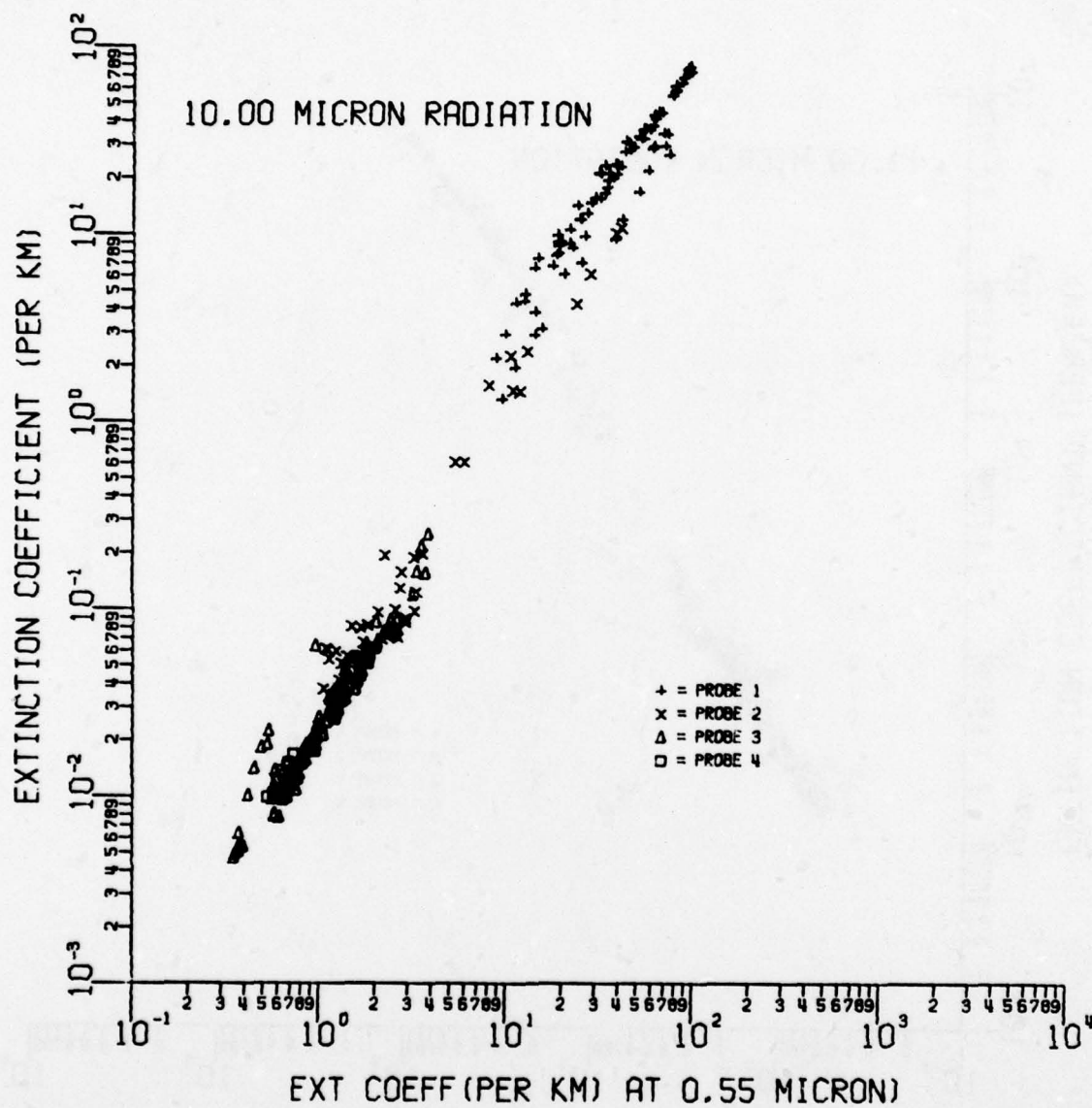


Figure A7-f

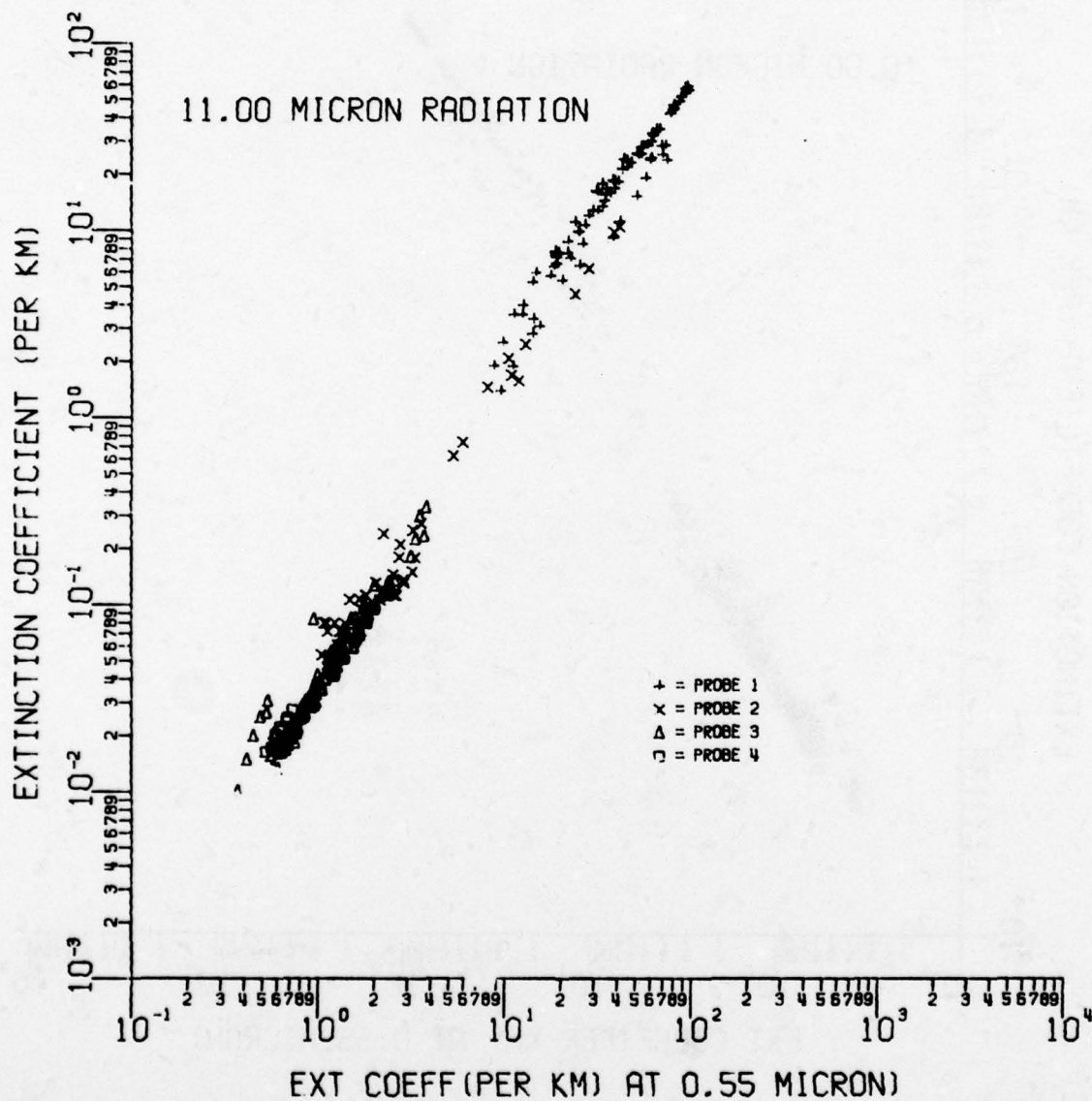


Figure A7-g

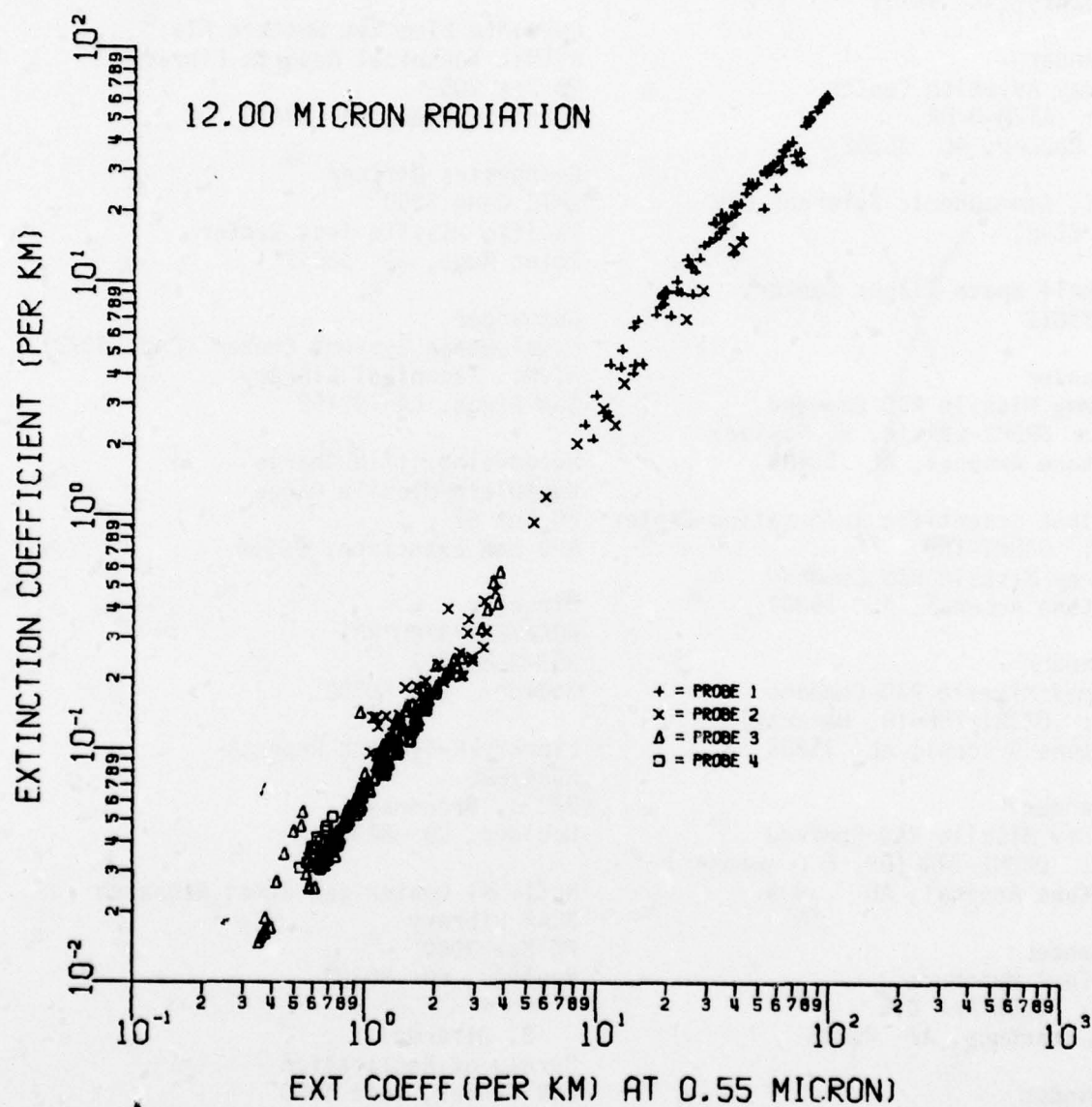


Figure A7-h

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